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OM protein - protein search, using sw model

Run on: May 4, 1999, 12:32:39 ; Search time 27.23 Seconds
(without alignments)
11.141 Million cell updates/sec

Title: US-09-037-460-2_COPY_55_69

Sequence: 78 1 RVCAAGRGTCYRTV 15

Scoring table: PAM150

Searched: 162890 seqs, 20225328 residues

Database: A_Geneseq_34:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	78	100.0	184	1 R98994	Vascular IBP-like
2	34	43.6	18	1 R57843	Vnrbeta3-7, bindin
3	33	42.3	18	1 R57842	Vnrbeta3-4, bindin
4	35	44.9	45	1 W10165	Alpha-hordothionin
5	35	44.9	50	1 W76687	Elapidae modified
6	36	46.2	75	1 R31601	Chicken nov protei
7	36	46.2	76	1 R31600	Chicken nov protei
8	35	44.9	60	1 W76646	Elapidae modified
9	35	44.9	67	1 W76685	Elapidae modified
10	35	44.9	70	1 W76685	Elapidae modified
11	35	44.9	72	1 W76684	Elapidae modified
12	35	44.9	73	1 W76684	Elapidae modified
13	35	44.9	74	1 W76661	Elapidae modified
14	30	38.5	11	1 R93364	MAGE-1 derived imm
15	31	39.7	24	1 W03034	Thrombolytic enzym
16	31	39.7	25	1 R22778	MVIIIB omega conoto
17	31	39.7	25	1 R37753	MVIIIB/SNX-159. Red
18	31	39.7	25	1 R39609	Omega conotoxin MV
19	31	39.7	25	1 R76090	Omega conopeptide
20	31	39.7	25	1 W12968	Natural omega-cono
21	31	39.7	25	1 W19545	Conus genus natura
22	31	39.7	25	1 W72606	Dendroides canad
23	30	38.5	18	1 R57849	New lipopolysaccha
24	31	39.7	18	1 W07695	Antiviral peptide
25	29	37.2	14	1 R47977	Tachyplesin I. Wat
26	29.5	37.8	17	1 P91671	Gigaslin II. Novel
27	29.5	37.8	17	1 R06266	Bacterial shock tr
28	29.5	37.8	17	1 R06266	Bacterial shock tr
29	29.5	37.8	17	1 R06266	Tachyplesin-II. Be
30	29.5	37.8	17	1 R08202	Tachyplesin-I. Beta
31	29.5	37.8	17	1 R23112	Tachyplesin-I. Beta
32	29.5	37.8	17	1 R23112	Tachyplesin-I. Beta
33	29.5	37.8	17	1 R23114	Tachyplesin-I. Beta
34	29.5	37.8	17	1 R38490	Tachyplesin-I. Beta
35	29.5	37.8	17	1 R38490	Tachyplesin-I. Beta
36	29.5	37.8	17	1 R38490	Tachyplesin-I. Beta
37	29.5	37.8	17	1 R38490	Tachyplesin-I. Beta
38	29.5	37.8	17	1 R38490	Tachyplesin-I. Beta
39	29.5	37.8	17	1 W66465	Cationic peptide t
40	28	35.9	10	1 W66465	Cationic peptide t
41	38	48.7	504	1 W20777	Targeting peptide
42	28	35.9	11	1 P91265	Haematopoietic ste
43	29	37.2	17	1 W78913	Tissue plasminogen
					Rat cocaine and am

44 28 35.9 12 1 W52102
45 29 37.2 18 1 R57840

ALIGNMENTS

RESULT 1
R98994
ID R98994 standard; Protein; 184 AA.
AC R98994;
DE 06-NOV-1996 (first entry)
KW Vascular IBP-like growth factor.
KW Vascular IBF-like growth factor; VIGF;
KW insulin-like growth factor binding protein; agonist; antagonist;
KW muscle wastage; osteoporosis; implant fixation; wound healing;
KW therapy; diagnosis.
OS Homo sapiens.
FH Key
FT Location/Qualifiers
FT 1..21
FT /label= sig_peptide
PN W09617931-A1.
PD 13-JUN-1996.
PF 09-DEC-1994; U14388.
PR 09-DEC-1994; WO-U14388.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Hastings GA, Rosen CA;
DR WPI; 96-287176/29.
DR N-PSDB; T34991.
PT Human vascular insulin-like growth factor binding protein-like
PT growth factor, and its nucleic acid sequence and (antagonists
PT used, e.g. to treat muscle wasting diseases or aid implant fixation,
PT or limit excess connective tissue prodn. during wound healing.
PS Claim 14; Page 43-44; 61pp; English.
CC Human vascular insulin-like growth factor binding protein-like
CC growth factor (R98994), or VIGF, is a protein of primarily
CC protein families. It can be expressed in e.g. E. coli, CHO or
CC insect host cells using a vector incorporating a cDNA clone
CC (T34991), or its derivative, obtd. from human umbilical
CC endothelial cells. It is useful therapeutically e.g. for
CC treating muscle wasting diseases or osteoporosis, or can be used
CC to detect diseases associated with under- or over-expression of VIGF,
CC or to screen for antagonists useful during wound healing.
SQ Sequence 184 AA;

Query Match 100.0%; Score 78; DB 1; Length 184;
Best Local Similarity 100.0%; Pred. No. 7e-05;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RVCAAGRGTCYRTV 15
Db 55 RVCAAGRGTCYRTV 69
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RESULT 2
R57843
ID R57843 standard; peptide; 18 AA.
AC R57843;
DE 28-MAR-1995 (first entry)
KW Vnrbeta3-7, binding site for vitronectin receptor alpha-v, beta-3.
KW Binding site; CDR; complementarity determining region; immunoglobulin;
KW heavy; light; primer extension; PCR; amplify; fibronectin; vitronectin;
KW RGD-dependent; integrin ligand; von Willebrand factor; EVB; gp350/220;
KW envelope glycoprotein; HIV; gp120; reovirus; hemagglutinin; insulin;
KW cellular receptor; CR2; CD4; hormone; thyroid stimulating hormone; TSH;
KW non-RGD-dependent; vitronectin; apo E; apo AI; MHC; class I; class II;
KW anti-gp120/IIIA; monoclonal antibody; alpha-v, beta-3; modulation;
KW coagulation; inflammation; anti-vitronectin; tumour cell adhesion;
KW migration.
OS Homo sapiens.

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PN WO9418221-A.
PD 18-AUG-1994.
PF 02-FEB-1994; U01258.
PR 02-FEB-1993; US-012566.
PR 28-JUN-1993; US-084542.
PA (SCRI ) SCRIPPS RES INST.
PI Barbas CF, Lerner RA;
DR WPI; 94-279675/34.
PT Production of binding sites within CDR regions of immunoglobulins
PT - displayed on the surface of filamentous phage particles, for
PT Inhibiting platelet aggregation and vitronectin binding
PS Claim 44; Page 22; 207pp; English.
CC The sequences given in R57837-84 are binding sites which were used in
CC the method of the invention for producing a polypeptide having a
CC binding site capable of binding a preselected agent. Nucleotide
CC sequences encoding these binding site peptides were introduced into
CC a CDR region of a nucleic acid encoding an immunoglobulin heavy (H)
CC or light (L) chain, by amplifying the CDR region by primer extension.
CC Preferred binding sites are derived from the RGD-dependent integrin
CC ligands, eg. fibronectin, vitronectin, von Willebrand factor, from
CC the envelope glycoprotein from viruses such as HIV gp120, EBV gp350/
CC CD4, from protein hormones such as thyroid stimulating hormone (TSH),
CC insulin, transferrin, from apolipoproteins such as apo E and apo AI,
CC from immunoglobulin CDRs and from MHC class I or II proteins. Non-RGD-
CC dependent integrin binding sites were selected for the affinity to bind
CC vitronectin receptor alpha-v, beta-3. An anti-gp11b/IIa monoclonal
CC antibody (Mab) produced in this way can be used to modulate platelet
CC adhesion in the treatment of coagulation and some inflammatory responses.
CC An anti-vitronectin Mab can be used in the treatment of cancer by
CC blocking tumour cell adhesion and migration. This sequence represents
CC an RGD-dependent binding site which has been shown to bind the human
CC vitronectin receptor (VnR) alpha-v, beta-3 when present in a phagemid
CC display protein.
SQ Sequence 18 AA;

Query Match 43.6%; Score 34; DB 1; Length 18;
Best Local Similarity 58.3%; Pred. No. 27;
Matches 7; Conservative 2; Mismatches 1; Indels 2; Gaps 1;

QY 3 CAAGRGET--CY 12
Dl : : : : :
Db 3 CTQGRGDFRNCY 14

RESULT 3
R57842
ID R57842 standard; peptide; 18 AA.
AC R57842;
DT 28-MAR-1995 (first entry)
DE VnRbeta3-4, binding site for vitronectin receptor alpha-v, beta-3.
KW Binding site; CDR; complementarity determining region; immunoglobulin;
KW heavy; light; primer extension; PCR; amplify; fibronectin; vitronectin;
KW RGD-dependent; integrin ligand; von Willebrand factor; EBV; gp350/220;
KW envelope glycoprotein; HIV; gp120; reovirus; hemagglutinin; insulin;
KW cellular receptor; CR2; CD4; hormone; thyroid stimulating hormone; TSH;
KW transferrin; apolipoprotein; apo E; apo AI; MHC; class I; class II;
KW non-RGD-dependent; vitronectin receptor; alpha-v, beta-3; modulation;
KW anti-gp11b/IIa; monoclonal antibody; Mab; platelet adhesion; cancer;
KW coagulation; inflammation; anti-vitronectin; tumour cell adhesion;
KW migration.
OS Homo sapiens.
PN WO9418221-A.
PD 18-AUG-1994.
PF 02-FEB-1994; U01258.
PR 02-FEB-1993; US-012566.
PR 28-JUN-1993; US-084542.
PA (SCRI ) SCRIPPS RES INST.
PI Barbas CF, Lerner RA;
DR WPI; 94-279675/34.
PT Production of binding sites within CDR regions of immunoglobulins
PT - displayed on the surface of filamentous phage particles, for
PT Inhibiting platelet aggregation and vitronectin binding
PS Claim 44; Page 22; 207pp; English.
CC The sequences given in R57837-84 are binding sites which were used in
CC the method of the invention for producing a polypeptide having a
CC binding site capable of binding a preselected agent. Nucleotide
CC sequences encoding these binding site peptides were introduced into
CC a CDR region of a nucleic acid encoding an immunoglobulin heavy (H)
CC or light (L) chain, by amplifying the CDR region by primer extension.
CC Preferred binding sites are derived from the RGD-dependent integrin
CC ligands, eg. fibronectin, vitronectin, von Willebrand factor, from
CC the envelope glycoprotein from viruses such as HIV gp120, EBV gp350/
CC CD4, from protein hormones such as thyroid stimulating hormone (TSH),
CC insulin, transferrin, from apolipoproteins such as apo E and apo AI,
CC from immunoglobulin CDRs and from MHC class I or II proteins. Non-RGD-
CC dependent integrin binding sites were selected for the affinity to bind
CC vitronectin receptor alpha-v, beta-3. An anti-gp11b/IIa monoclonal
CC antibody (Mab) produced in this way can be used to modulate platelet
CC adhesion in the treatment of coagulation and some inflammatory responses.
CC An anti-vitronectin Mab can be used in the treatment of cancer by
CC blocking tumour cell adhesion and migration. This sequence represents
CC an RGD-dependent binding site which has been shown to bind the human
CC vitronectin receptor (VnR) alpha-v, beta-3 when present in a phagemid
CC display protein.
SQ Sequence 18 AA;

Query Match 42.3%; Score 33; DB 1; Length 18;
Best Local Similarity 50.0%; Pred. No. 38;
Matches 6; Conservative 3; Mismatches 1; Indels 2; Gaps 1;

QY 3 CAAGRGE--TCY 12
Dl : : : : :
Db 3 CTQGRGDFRNCY 14

RESULT 4
W10165
ID W10165 standard; protein; 45 AA.
AC W10165;
DT 15-JUL-1997 (first entry)
DE Alpha-hordothionin threonine rich amino acid sequence.
KW Animal feed; food; barley.
KW Hordeum vulgare.
OS Synthetic.
PN WO9638562-A1.
PD 05-DEC-1996.
PR 31-MAY-1996; U08219.
PR 02-JUN-1995; US-459180.
PA (PION-) PIONEER HI-BRED INT INC.
PI Rao GA;
DR WPI; 97-034375/03.
PT New modified alpha-hordothionin having threonine amino acid substns.
PT - to increase the threonine content of e.g. animal feed
PS Claim 1; Page 11; 19pp; English.
CC The present sequence is a threonine rich alpha-hordothionin amino
CC acid sequence. The protein contains a threonine residue at positions
CC 1, 5, 7, 8, 11, 15, 17, 19, 22, 23, 24, 30, 32, 34, 38, and 41. The
CC threonine has substituted polar, charged and hydrophobic residues.
CC With exception of arginine at position 10, serine at position 2 and
CC lysine at position 45, as they are required for maintaining the structure
CC of the protein through a hydrogen-bonding network. The protein produced
CC can be used in foods or feeds to provide higher levels of essential amino
CC acid threonine.
SQ Sequence 45 AA;

Query Match 44.9%; Score 35; DB 1; Length 45;
Best Local Similarity 50.0%; Pred. No. 43;
Matches 6; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 3 CAAGRGETCYRT 14
Dl : : : : :

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CC or multifunctional activities against blood coagulation, particularly
 CC thrombus formation and arterial/venous wall thickening at the sites of
 CC injury. The variants may have activities against leukocyte recruitment,
 CC immune system activation, tissue fibrosis and tumorigenesis. The
 CC polypeptides can be used for the treatment or prophylaxis of a disease
 CC associated with thrombosis, e.g. myocardial infarction, retinal
 CC neovascularisation, endothelial injury, dysregulated apoptosis, abnormal
 CC cell migration, leukocyte recruitment, immune system activation, tissue
 CC fibrosis or tumorigenesis.
 SQ Sequence 70 AA;

Query Match 44.9%; Score 35; DB 1; Length 70;
 Best Local Similarity 22.2%; Pred. No. 63;
 Matches 6; Conservative 6; Mismatches 3; Indels 12; Gaps 1;
 QY 1 RYCAARG-----EFCYRTV 15
 DB 1 RICFTPRGDMPGYPGQEDSCYKNI 27

RESULT 11

W76684

ID W76684 standard; Protein: 72 AA.

AC W76684;

DT 05-JAN-1999 (first entry)

DE Elapidae modified dendroaspin protein fragment Den-Hr.

KW Dendroaspin; snake venom; clotting cascade; anticoagulant; platelet;

KW integrin binding; injury; blood; cell migration; thrombosis; inhibitor;

KW proliferation; signal transduction; regulator; coagulation; treatment;

KW prophylactic; artery; vein; wall thickening; myocardial infarction;

KW retinal neovascularisation; dysregulated apoptosis; tumorigenesis;

KW leukocyte recruitment, immune system; tissue fibrosis.

OS Elapidae.

OS Synthetic.

PN W09842834-Al.

PD 01-OCT-1998.

PF 20-MAR-1998; G00848.

PR 20-MAR-1997; GB-005787.

PA (THRO-) THROMBOSIS RES. INST.

PI Authi K, Kakkar V, Lu X, Scully MF;

DR WPI; 98-542278/46.

PT New hybrid dendroaspin polypeptide(s) - used for treating, e.g.

PT thrombosis, myocardial infarction, dysregulated apoptosis, abnormal

PT cell migration and immune system activation

PS Claim 6; Fig 3C; 59pp; English.

CC W76645-W76688 represent modified dendroaspin protein fragments isolated

CC from snake venom. When dendroaspin is modified to incorporate further

CC functional amino acid sequence, e.g. active portions or motifs of

CC agonists, antagonists or inhibitors of factors in the clotting cascade,

CC the resulting molecules are particularly useful as anticoagulants. The

CC molecules have an integrin binding activity which when administered in

CC vivo results in the binding of the platelets at sites of injury. Non-wild

CC inhibiting the aggregation of the platelets at sites of injury. Non-wild

CC type dendroaspin domains provide secondary, optionally further

CC proliferation and regulating signal transduction, inhibiting cell migration and

CC or multifunctional activities against blood coagulation. Such variants have bi-

CC thrombus formation and arterial/venous wall thickening at the sites of

CC injury. The variants may have activities against leukocyte recruitment,

CC immune system activation, tissue fibrosis and tumorigenesis. The

CC polypeptides can be used for the treatment or prophylaxis of a disease

CC associated with thrombosis, e.g. myocardial infarction, retinal

CC neovascularisation, endothelial injury, dysregulated apoptosis, abnormal

CC cell migration, leukocyte recruitment, immune system activation, tissue

CC fibrosis or tumorigenesis.
 SQ Sequence 72 AA;

Query Match 44.9%; Score 35; DB 1; Length 72;
 Best Local Similarity 22.2%; Pred. No. 65;
 Matches 6; Conservative 6; Mismatches 3; Indels 12; Gaps 1;

QY 1 RYCAARG-----EFCYRTV 15
 DB 1 RICFTPRGDMPGYPGQEDSCYKNI 27

RESULT 12

W76658

ID W76658 standard; Protein: 73 AA.

AC W76658;

DT 05-JAN-1999 (first entry)

DE Elapidae modified dendroaspin protein fragment DEN-HR11.

KW Dendroaspin; snake venom; clotting cascade; anticoagulant; platelet;

KW integrin binding; injury; blood; cell migration; thrombosis; inhibitor;

KW proliferation; signal transduction; regulator; coagulation; treatment;

KW prophylactic; artery; vein; wall thickening; myocardial infarction;

KW retinal neovascularisation; dysregulated apoptosis; tumorigenesis;

KW leukocyte recruitment, immune system; tissue fibrosis.

OS Elapidae.

OS Synthetic.

PN W09842834-Al.

PD 01-OCT-1998.

PF 20-MAR-1998; G00848.

PR 20-MAR-1997; GB-005787.

PA (THRO-) THROMBOSIS RES. INST.

PI Authi K, Kakkar V, Lu X, Scully MF;

DR WPI; 98-542278/46.

PT New hybrid dendroaspin polypeptide(s) - used for treating, e.g.

PT thrombosis, myocardial infarction, dysregulated apoptosis, abnormal

PT cell migration and immune system activation

PS Claim 6; Fig 3A; 59pp; English.

CC W76645-W76688 represent modified dendroaspin protein fragments isolated

CC from snake venom. When dendroaspin is modified to incorporate further

CC functional amino acid sequence, e.g. active portions or motifs of

CC agonists, antagonists or inhibitors of factors in the clotting cascade,

CC the resulting molecules are particularly useful as anticoagulants. The

CC molecules have an integrin binding activity which when administered in

CC vivo results in the binding of the platelets at sites of injury. Non-wild

CC inhibiting the aggregation of the platelets at sites of injury. Non-wild

CC type dendroaspin domains provide secondary, optionally further

CC proliferation and regulating signal transduction, inhibiting cell migration and

CC or multifunctional activities against blood coagulation. Such variants have bi-

CC thrombus formation and arterial/venous wall thickening at the sites of

CC injury. The variants may have activities against leukocyte recruitment,

CC immune system activation, tissue fibrosis and tumorigenesis. The

CC polypeptides can be used for the treatment or prophylaxis of a disease

CC associated with thrombosis, e.g. myocardial infarction, retinal

CC neovascularisation, endothelial injury, dysregulated apoptosis, abnormal

CC cell migration, leukocyte recruitment, immune system activation, tissue

CC fibrosis or tumorigenesis.
 SQ Sequence 73 AA;

Query Match 44.9%; Score 35; DB 1; Length 73;
 Best Local Similarity 22.2%; Pred. No. 66;
 Matches 6; Conservative 6; Mismatches 3; Indels 12; Gaps 1;

QY 1 RYCAARG-----EFCYRTV 15
 DB 1 RICFTPRGDMPGYPGQEDSCYKNI 27

RESULT 13

W76661

ID W76661 standard; Protein: 74 AA.

AC W76661;

DT 05-JAN-1999 (first entry)

DE Elapidae modified dendroaspin protein fragment DEN-TM11.

KW Dendroaspin; snake venom; clotting cascade; anticoagulant; platelet;

KW integrin binding; injury; blood; cell migration; thrombosis; inhibitor;

KW proliferation; signal transduction; regulator; coagulation; treatment;

KW prophylactic; artery; vein; wall thickening; myocardial infarction;

KW retinal neovascularisation; dysregulated apoptosis; tumorigenesis;

KW leukocyte recruitment, immune system; tissue fibrosis.
OS Elapidae.
OS Synthetic.
PN WO9842834-A1.
PD 01-OCT-1998.
PF 20-MAR-1998; G00848.
PR 20-MAR-1997; GB-005787.
PA (THRO-) THROMBOSIS RES INST.
PI Authi K, Kakkar V, Lu X, Scully MF;
DR WPI: 98-542278/46.
PT New hybrid dendroaspin polypeptide(s) - used for treating, e.g.
PT thrombosis, myocardial infarction, dysregulated apoptosis, abnormal
PT cell migration and immune system activation
PS Claim 6; Fig 3A; 59pp; English.
CC W76645-W76688 represent modified dendroaspin protein fragments isolated
CC from snake venom. When dendroaspin is modified to incorporate further
CC functional amino acid sequence, e.g. active portions or motifs of
CC agonists, antagonists or inhibitors of factors in the clotting cascade,
CC the resulting molecules are particularly useful as anticoagulants. The
CC molecules have an integrin binding activity which when administered in
CC vivo results in the binding of the molecules to platelets thereby
CC inhibiting the aggregation of the platelets at sites of injury. Non-wild
CC type dendroaspin domains provide secondary, optionally further
CC functionality, e.g. antithrombotic action, inhibiting cell migration and
CC proliferation and regulating signal transduction. Such variants have bi-
CC or multifunctional activities against blood coagulation, particularly
CC thrombus formation and arterial/venous wall thickening at the sites of
CC injury. The variants may have activities against leukocyte recruitment,
CC immune system activation, tissue fibrosis and tumorigenesis. The
CC polypeptides can be used for the treatment or prophylaxis of a disease
CC associated with thrombosis, e.g. myocardial infarction, retinal
CC neovascularisation, endothelial injury, dysregulated apoptosis, abnormal
CC cell migration, leukocyte recruitment, immune system activation, tissue
CC fibrosis or tumorigenesis.
SQ Sequence 74 AA;

Query Match 44.9%; Score 35; DB 1; Length 74;
Best Local Similarity 22.2%; Pred. No. 66;

Matches 6; Conservative 6; Mismatches 3; Indels 12; Gaps 1;

QY 1 RVCAGRG-----ETCYRTV 15
I : I : I :
D 1 RICPTRGDPGPPGPGQEDSCYKNI 27

RESULT 14
R89364

ID R89364 standard; peptide; 11 AA.

AC R89364;
DE 18-SEP-1996 (first entry)
DE MAGE-1 derived immunogenic peptide.
KW Immunogenic peptide; supermotif; HLA molecule; CTL response;
KW therapeutic; diagnostic; cancer; viral infection; hepatitis B;
KW hepatitis C.
OS Synthetic.

PN WO9603140-A1.
PD 08-FEB-1996.

PF 21-JUL-1995; U09234.

PR 21-JUL-1994; US-278634.

PR 23-NOV-1994; US-344824.

PR 30-MAY-1995; US-452843.

PA (CYTE-) CYTEL CORP.

PI Sette A, Sidney J;

DR WPI: 96-116784/12.

PT Compn. comprising immunogenic peptide with supermotif allowing more
PT than one HLA mol. to bind - used to induce CTL response in patient
PT and for in vivo and ex vivo therapeutic and diagnostic applications
PS Claim 2; Page 26; 32pp; English.
CC The sequences given in R89362-82 are immunogenic peptides which were
CC use in the composition of the invention. The composition comprises
CC an immunogenic peptide of 9-10 residues with a supermotif which
CC allows binding of more than one HLA molecule. It pref. comprises

CC two conserved residues, a first at the 2nd position from the N-
CC terminal is Pro, and a 2nd at the C-terminal is Met. These peptides
CC are used to induce a CTL response in a patient. They are also
CC useful in compositions for in vivo and ex vivo therapeutic and
CC diagnostic applications, e.g. the treatment of cancer and viral
CC infections, e.g. hepatitis B and C.
SQ Sequence 11 AA;

Query Match 38.5%; Score 30; DB 1; Length 11;
Best Local Similarity 57.1%; Pred. No. 68;
Matches 4; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 9 ETCYRTV 15
I : I : I :
D 4 ESCFRAV 10

RESULT 15
W03034

ID W03034 standard; peptide; 24 AA.

AC W03034;

DT 17-FEB-1997 (first entry)

DE Thrombolytic enzyme fragment #3.

KW Thrombolytic enzyme; destabilase; hydrolysis; D fragment dimer;

KW epsilon-(gamma-glutamyl)-lysine isopeptide bond; stabilised fibrin;

KW salivary gland; medicinal leech.

OS Hirudo medicinalis.

PN WO9619999-A1.

PD 04-JUL-1996.

PF 25-MAY-1995; RU0102.

PR 23-DEC-1994; RU-043698.

PA (BIOO-) INST BIOORG KHIM IM M SHERY.

PI Barsova EV, Baskova IP, Bogdanova EA, Lukyanov SA;

PI Sverdllov ED, Zavalova IL;

DR WPI: 96-321638/32.

PT Thrombolytic enzyme destabilase - selectively hydrolysing stabilised
PT fibrin

PS Disclosure; Page 3; 16pp; Russian

CC The sequences given in W03032-35 represent CNBr fragments of the novel
CC thrombolytic enzyme of the invention. The thrombolytic enzyme, also
CC known as destabilase, is capable of hydrolysing exo- and endo-

CC epsilon-(gamma-glutamyl)-lysine isopeptide bonds in synthetic

CC substrates and in stabilised fibrin and its D fragment dimer.

CC Destabilase may be used as a thrombolytic agent in clinical and

CC experimental medicine. It can be isolated from the salivary glands

CC of medicinal leeches.

SQ Sequence 24 AA;

Query Match 39.7%; Score 31; DB 1; Length 24;
Best Local Similarity 55.6%; Pred. No. 96;

Matches 5; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 3 CAAGRGTC 11

D 7 CTGGROPTC 15

Search completed: May 4, 1999, 12:32:41
Job time: 9788 sec

GenCore version 4.5
Copyright (c) 1993 - 1998 Compugen Ltd.
OM protein - protein search, using sw model
Run On: May 4, 1999, 08:18:15 ; Search time 25.07 Seconds
(without alignments)
22.413 Million cell updates/sec

Title: US-09-037-460-2_COPY_30_44
Perfect score: 74
Sequence: 1 QHCDSSCKSPRCK 15

Scoring table: PAM150

Searched: 116738 seqs, 37460341 residues

Database : PIR_58.*

- 1: PIR1.*
- 2: PIR2.*
- 3: PIR3.*
- 4: PIR4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	40	54.1	57	2 S59073	metallothionein is
2	42	56.8	157	2 S39849	PfrrA protein - Pse
3	44	59.5	861	2 A48825	Notch homolog Motc
4	42	56.8	488	1 EXHU	coagulation factor
5	38	51.4	75	2 S17156	metallothionein -
6	40	54.1	251	2 A5035	cysteine-rich prot
7	37	50.0	61	2 A5049	metallothionein pr
8	36	48.6	39	2 A46057	thrombin inhibitor
9	41	55.4	473	2 A56175	adhesive plaque pr
10	42	56.8	772	2 S32659	integrin beta 2 ch
11	36	48.6	57	1 SKWDS	metallothionein 2
12	44	59.5	2524	2 A35844	Kotch protein - Af
13	44	59.5	2531	2 A46019	gene Notch-1 prote
14	37	50.0	111	2 S44787	D2007.1 protein -
15	36	48.6	77	2 S36032	thrombin inhibitor
16	39	52.7	400	2 S32804	beta-3-adrenergic
17	39	52.7	400	2 A41679	beta-3-adrenergic
18	39	52.7	400	2 A53281	beta-3-adrenergic
19	37.5	50.7	200	2 S35292	hypothetical prote
20	39	52.7	410	2 S15163	probable transpos
21	37	50.0	180	2 A45810	glycoprotein anti
22	38	51.4	319	2 A53502	folistatin - Afri
23	35	47.3	79	2 H69193	2-oxoisovalerate o
24	38	51.4	335	2 S71796	centrosome-binding
25	38	51.4	343	2 S55369	folistatin - chlc
26	38	51.4	379	1 DEILSP	alcohol dehydrogen
27	38	51.4	379	1 S01893	alcohol dehydrogen
28	42	56.8	2555	2 A40043	notch protein homo
29	36	48.6	155	2 A45293	conopressin precu
30	39.5	53.4	837	2 A42112	mucin-like peptide
31	38	51.4	414	2 S36838	acetyl-CoA C-acyt
32	39	52.7	683	2 JC5393	zinc finger protei
33	38	51.4	466	1 KFBU7	coagulation factor
34	34	45.9	75	2 B45206	metallothionein 2
35	35	47.3	125	1 NFRT1	oxytocin / neuroph
36	35	47.3	125	1 A43755	oxytocin / neuroph
37	34	45.9	78	2 S09414	proteinase inhibit
38	40	54.1	1375	2 F48216	neurexin iii-alpha
39	40	54.1	1378	2 E48216	neurexin iii-alpha

ALIGNMENTS

RESULT 1

S59073

metallothionein isoform Iia - blue crab

C:Species: Callinectes sapidus

C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 29-Aug-1997

C:Accession: S59073

R:Brouwer, M.; Enghild, J.; Hoexum-Brouwer, T.; Thogersen, I.; Truncali, A.

Biochem. J. 311, 617-622, 1995

A:Title: Primary structure and tissue-specific expression of blue crab (Callinectes s

A:Reference number: S59072

A:Accession: S59073

A:Molecule type: protein

A:Residues: 1-57 <BRO>

C:Superfamily: metallothionein

C:Keywords: metal binding

Query Match 54.1%; Score 40; DB 2; Length 57;

Best Local Similarity 46.2%; Pred. No. 7;

Matches 6; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 3 CDSSECKSPRCK 15

DB 11 CREGECKTGCKCK 23

RESULT 2

S39849

PfrrA protein - Pseudomonas putida

C:Species: Pseudomonas putida

C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 01-Aug-1997

C:Accession: S39849

R:Venturi, V.; Ottavanger, C.; Leong, J.; Weisbeek, P.J.

Mol. Microbiol. 10, 63-73, 1993

A:Title: Identification and characterization of a siderophore regulatory gene (pfra)

A:Reference number: S39849

A:Accession: S39849

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-157 <VEN>

Query Match

Best Local Similarity 56.8%; Score 42; DB 2; Length 157;

Matches 6; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 QHCDSSCKSPR 13

DB 107 DHCEKGECKDPER 119

RESULT 3

A48825

Notch homolog Motch protein - mouse (fragment)

C:Species: Mus musculus (house mouse)

C:Date: 01-Dec-1993 #sequence_revision 18-Nov-1994 #text_change 14-Aug-1998

C:Accession: A48825

R:Reaume, A.G.; Conlon, R.A.; Zirngibl, R.; Yamaguchi, T.P.; Rossant, J.

Dev. Biol. 154, 377-387, 1992

A:Title: Expression analysis of a Notch homologue in the mouse embryo.

A:Reference number: A48825; MUID:93050801

A:Accession: A48825

A:Status: preliminary: not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-861 <REA>
A:Experimental source: embryo
A:Note: sequence extracted from NCBI backbone (NCBIP:119144)
C:Superfamily: unassigned ankryin repeat proteins; ankryin repeat homology; EGF homology
F:26-57/Domain: EGF homology <EGF>

Query Match 59.5%; Score 44; DB 2; Length 861;
Best Local Similarity 58.3%; Pred. No. 12;
Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 3 CDSSECKSSPRCK 14
||||: |||: ||
DB 274 CDSAPCKNGGRC 285

RESULT 4
EXHU
coagulation factor Xa (EC 3.4.21.6) precursor - human
N:Alternate names: Stuart factor
C:Species: Homo sapiens (man)
C:Date: 15-Nov-1984 #sequence_revision 02-May-1994 #text_change 24-Oct-1997
C:Accession: A24478; JQ0917; A42485; A25853; A22208; A21284; A20362; S39415; I54051; A00
R:Leytus, S.P.; Foster, D.C.; Kurachi, K.; Davie, E.W.
Biochemistry 25, 5098-5102, 1986
A:Title: Gene for human factor X: a blood coagulation factor whose gene organization is
A:Reference number: A24478; MUID:87026600
A:Accession: A24478
A:Molecule type: DNA
A:Residues: 1-488 <LEY>
A:Cross-references: GB:L29433; GB:M14327; NID:9459809; PID:g182831
R:Messier, T.L.; Pittman, D.D.; Long, G.L.; Kaufman, R.J.; Church, W.R.
Gene 99, 291-294, 1991
A:Title: Cloning and expression in COS-1 cells of a full-length cDNA encoding human coag
A:Reference number: JQ0917; MUID:91216473
A:Accession: JQ0917
A:Molecule type: mRNA
A:Residues: 1-488 <MES>
A:Cross-references: GB:M57285; NID:g182389; PID:g182390
R:Miao, C.H.; Leytus, S.P.; Chung, D.W.; Davie, E.W.
J. Biol. Chem. 267, 7395-7401, 1992
A:Title: Liver-specific expression of the gene coding for human factor X, a blood coagul
A:Reference number: A42485; MUID:92218390
A:Accession: A42485
A:Molecule type: DNA
A:Residues: 1-15 <MTA>
A:Experimental source: liver
A:Note: sequence extracted from NCBI backbone (NCBIN:93780, NCBIP:93787)
R:Kaul, R.K.; Hildebrand, B.; Roberts, S.; Jagadeeswaran, P.
Gene 41, 311-314, 1986
A:Title: Isolation and characterization of human blood-coagulation factor X cDNA.
A:Reference number: A25853; MUID:86221713
A:Accession: A25853
A:Molecule type: mRNA
A:Residues: 19-284, 'E', 289-488 <KAU>
A:Cross-references: GB:M2613; NID:g180335; PID:g180336
R:Fung, M.R.; Hay, C.W.; MacGillivray, R.T.A.
Proc. Natl. Acad. Sci. U.S.A. 82, 3591-3595, 1985
A:Title: Characterization of an almost full-length cDNA coding for human blood coagulation
A:Reference number: A22208; MUID:85216545
A:Accession: A22208
A:Molecule type: mRNA
A:Residues: 13-441, 'S', 443-488 <FUN>
A:Cross-references: GB:K03194; NID:g182840; PID:g182841
R:Leytus, S.P.; Chung, D.W.; Kistiel, W.; Kurachi, K.; Davie, E.W.
Proc. Natl. Acad. Sci. U.S.A. 81, 3699-3702, 1984
A:Title: Characterization of a cDNA coding for human factor X.
A:Reference number: A21284; MUID:84222026
A:Accession: A21284
A:Molecule type: mRNA
A:Residues: 13-284, 'E', 289-488 <LE2>

A:Cross-references: GB:K01886
R:McMullen, B.A.; Fujikawa, K.; Kistiel, W.; Sasagawa, T.; Howald, W.N.; Kwa, E.Y.; We
Biochemistry 22, 2875-2884, 1983
A:Title: Complete amino acid sequence of the light chain of human blood coagulation f
A:Reference number: A20362; MUID:83257207
A:Accession: A20362
A:Molecule type: protein
A:Residues: 41-179 <CMC>
R:Inoue, K.; Morita, T.
Eur. J. Biochem. 218, 153-163, 1993
A:Title: Identification of O-linked oligosaccharide chains in the activation peptides
A:Reference number: S39414
A:Accession: S39415
A:Molecule type: protein
A:Residues: 183-234 <INO>
A:Note: glycosylation sites
R:Jagadeeswaran, P.; Reddy, S.V.; Rao, K.J.; Hamsabhusanam, K.; Lyman, G.
Gene 84, 517-519, 1989
A:Title: Cloning and characterization of the 5' end (exon 1) of the gene encoding hum
A:Reference number: I54051; MUID:90128299
A:Accession: I54051
A:Status: translation not shown; translated from GB/EMBL/DDBJ
A:Molecule type: DNA
A:Residues: 1-23 <RES>
A:Cross-references: GB:M33297; NID:g183860; PID:G553330
R:Padmanabhan, K.; Padmanabhan, K.P.; Tullinsky, A.; Park, C.H.; Bode, W.; Huber, R.;
J. Mol. Biol. 232, 947-966, 1993
A:Title: Structure of human des(1-45) factor Xa at 2.2 angstroms resolution.
A:Reference number: A49458
A:Contents: annotation; X-ray crystallography, 2.2 angstroms
C:Comment: The two chains held together by one disulfide bond are formed from a singl
C:Comment: The activation peptide is cleaved by factor IXa (in the intrinsic pathway)
C:Genetics:
A:Gene: GDB:F10
A:Cross-references: GDB:119890; OMIM:227600
A:Map position: 13q34-13q34
A:Introns: 24/1; 77/3; 86/1; 124/1; 150/3; 249/3; 289/1
A:Note: deficiency of this factor causes Stuart disease
C:Function:
A:Description: catalyzes the proteolytic activation of prothrombin to thrombin in the
A:Pathway: blood coagulation
C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homol
C:Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxyglu
F:1-23/Domain: signal sequence #status predicted <SIG>
F:24-40/Domain: propeptide #status predicted <PRO>
F:25-84/Domain: Gla domain homology <GLA>
F:41-179/Product: coagulation factor X light chain #status experimental <LCH>
F:129-164/Domain: EGF homology <EGF1>
F:183-488/Product: coagulation factor X heavy chain #status experimental <HCH>
F:183-234/Domain: activation peptide #status experimental <APT>
F:235-488/Product: coagulation factor Xa heavy chain #status experimental <ACT>
F:235-462/Domain: trypsin homology <TRY>
F:46-47,54,56,59,60,65,66,69,72,79/Modified site: gamma-carboxyglutamic acid (Glu) #s
F:57-62/Disulfide bonds: #status predicted
F:90-101,95-110,112-121,129-140,136-149,151-164,172-342,241-246,261-277,390-404,415-4
F:103/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status experimental
F:199,211/Binding site: carbohydrate (Thr) (covalent) #status experimental
F:221,231/Binding site: carbohydrate (Asn) (covalent) #status experimental
F:234-235/Cleavage site: Arg-Ile (coagulation factor IXa, coagulation factor VIIa) #s
F:276,322,419/Active site: His, Asp, Ser #status experimental

Query Match 56.8%; Score 42; DB 1; Length 488;
Best Local Similarity 33.3%; Pred. No. 16;
Matches 5; Conservative 6; Mismatches 4; Indels 0; Gaps 0;

QY 1 QHCDSECKSSPRCK 15
||||: |||: ||
DB 88 DQCETSPQNGKCK 102

RESULT 5

S17156
 metallothionein - eastern oyster
 C:Species: Crassostrea virginica (eastern oyster)
 C:Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #text_change 08-Sep-1997
 C:Accession: S17156
 R:Unger, M.E.; Chen, T.T.; Murphy, C.M.; Vestling, M.M.; Fenselau, C.; Roeslajadi, G.
 Biochim. Biophys. Acta 1074, 371-377, 1991
 A:Title: Primary structure of molluscan metallothioneins deduced from PCR-amplified cDNA
 A:Reference number: S17156; MUID:91363394
 A:Accession: S17156
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-75 <UNG>
 A:Cross-references: EMBL:X59862; NID:g288277; PID:g288278
 C:Superfamily: metallothionein

Query Match 51.4%; Score 38; DB 2; Length 75;
 Best Local Similarity 46.2%; Pred. No. 16;
 Matches 6; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

Qy 3 CDSSECKSSPRCK 15

Db 19 CPATCKCGPGCK 31

RESULT 6

A55035
 cysteine-rich protein CRP1 - earthworm (Enchytraeus buchholzi)
 C:Species: Enchytraeus buchholzi
 C:Date: 14-Nov-1994 #sequence_revision 03-Nov-1995 #text_change 10-Sep-1997
 C:Accession: A55035; S45034
 R:Willuhn, J.; Schmitt-Wrede, H.P.; Greven, H.; Wunderlich, F.
 J. Biol. Chem. 269, 24688-24691, 1994
 A:Title: cDNA cloning of a cadmium-inducible mRNA encoding a novel cysteine-rich, non-me
 A:Reference number: A55035
 A:Accession: A55035
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-251 <WIL>
 A:Cross-references: EMBL:X79344; NID:g488802; PID:g488803

Query Match 54.1%; Score 40; DB 2; Length 251;
 Best Local Similarity 38.5%; Pred. No. 20;
 Matches 5; Conservative 5; Mismatches 3; Indels 0; Gaps 0;

Qy 3 CDSSECKSSPRCK 15

Db 223 CDNVNCKGSSCR 235

RESULT 7

S14049
 metallothionein precursor - yeast (Saccharomyces cerevisiae)
 A:Alternate names: protein YHR053c; protein YHR055c
 C:Species: Saccharomyces cerevisiae
 C:Date: 16-Sep-1992 #sequence_revision 16-Sep-1992 #text_change 05-Dec-1997
 C:Accession: S14049; A29373; A17610; A24123; S46703; S46705
 R:Jeyaprakash, A.; Welch, J.W.; Fogel, S.
 Mol. Gen. Genet. 225, 363-368, 1991
 A:Title: Multicopy CUP1 plasmids enhance cadmium and copper resistance levels in yeast.
 A:Reference number: S14049; MUID:91203809
 A:Accession: S14049
 A:Molecule type: DNA
 A:Residues: 1-61 <MOL>
 R:Karim, M.; Najarian, R.; Haslinger, A.; Valenzuela, P.; Welch, J.; Fogel, S.
 Proc. Natl. Acad. Sci. U.S.A. 81, 337-341, 1984
 A:Title: Primary structure and transcription of an amplified genetic locus: the CUP1 loc
 A:Reference number: A29373; MUID:84119482
 A:Accession: A29373
 A:Molecule type: DNA

A:Residues: 1-61 <KAR>
 A:Cross-references: EMBL:K02204; NID:g171337; PID:g171338
 R:Butt, T.R.; Sternberg, E.J.; Gorman, J.A.; Clark, P.; Hamer, D.; Rosenberg, M.; Cro
 Proc. Natl. Acad. Sci. U.S.A. 81, 3332-3336, 1984
 A:Title: Copper metallothionein of yeast, structure of the gene, and regulation of ex
 A:Reference number: A17610; MUID:84221953
 A:Accession: A17610
 A:Molecule type: DNA
 A:Residues: 1-61 <BUT>
 A:Cross-references: EMBL:K02204; NID:g171337; PID:g171338
 R:Winge, D.R.; Nielson, K.B.; Gray, W.R.; Hamer, D.H.
 J. Biol. Chem. 260, 14464-14470, 1985
 A:Title: Yeast metallothionein. Sequence and metal-binding properties.
 A:Reference number: A92506; MUID:86033949
 A:Accession: A24123
 A:Molecule type: protein
 A:Residues: 9-61 <WIN>
 R:Latreille, P.
 submitted to the EMBL Data Library, May 1994
 A:Description: The sequence of S. cerevisiae cosmid 8025.
 A:Reference number: S46691
 A:Accession: S46703
 A:Molecule type: DNA
 A:Residues: 1-61 <LAT>
 A:Cross-references: EMBL:U00061; NID:g487943; PID:el08807; MIPS:YHR053c
 A:Genetics: CUP1A
 A:Accession: S46705
 A:Molecule type: DNA
 A:Residues: 1-61 <LA2>
 A:Cross-references: EMBL:U00061; NID:g487943; PID:g487953; MIPS:YHR055c
 A:Genetics: CUP1B
 C:Genetics: <CUP1A>
 A:Gene: SGD:CUP1; CUP1A
 A:Cross-references: MIPS:YHR053c; SGD:S0001095
 A:Map position: 8R
 C:Genetics: <CUP1B>
 A:Gene: SGD:CUP1; CUP1B
 A:Cross-references: MIPS:YHR055c; SGD:S0001097
 A:Map position: 8R
 C:Superfamily: metallothionein

Query Match 50.0%; Score 37; DB 2; Length 61;
 Best Local Similarity 30.8%; Pred. No. 20;
 Matches 4; Conservative 6; Mismatches 3; Indels 0; Gaps 0;

Qy 2 HCDSECKSSPRC 14

Db 16 QCQCGCKNNEQC 28

RESULT 8

A46057
 thrombin inhibitor hemadin - terrestrial leech (Haemadipsa sylvestrus) (fragment)
 C:Species: Haemadipsa sylvestrus
 C:Date: 21-Sep-1993 #sequence_revision 18-Nov-1994 #text_change 20-Mar-1998
 C:Accession: A46057
 R:Strube, K.H.; Kroger, B.; Bialojan, S.; Otte, M.; Dödt, J.
 J. Biol. Chem. 268, 8590-8595, 1993
 A:Title: Isolation, sequence analysis, and cloning of haemadin. An anticoagulant pept
 A:Reference number: A46057; MUID:9332009
 A:Accession: A46057
 A:Status: preliminary
 A:Molecule type: mRNA; protein
 A:Residues: 1-39 <STR>
 A:Cross-references: GB:S58792; NID:g302486; PID:g302487
 A:Note: sequence extracted from NCBI backbone (NCBIN:129610, NCBIP:129611)

Query Match 48.6%; Score 36; DB 2; Length 39;
 Best Local Similarity 30.8%; Pred. No. 20;
 Matches 4; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

QY 1 QHCDSECKSSPR 13
| : : : : :
Db 12 QSCNDGQCSGDPK 24

RESULT 9
A56175
adhesive plaque protein Mgfp2 precursor - Mediterranean mussel
C:Species: Mytilus galloprovincialis (Mediterranean mussel)
C>Date: 27-Apr-1995 #sequence_revision 03-Oct-1995 #text_change 10-Sep-1997
R:Inoue, K.; Takeuchi, Y.; Miki, D.; Odo, S.
J. Biol. Chem. 270, 6698-6701, 1995
A:Title: Mussel adhesive plaque protein gene is a novel member of epidermal growth factor
A:Reference number: A56175
A:Accession: A56175
A:Molecule type: mRNA
A:Residues: 1-473 <INO>
A:Cross-references: GB:D43794; NID:g602767; PID:di008438; PID:g602768
C:Keywords: duplication
F:1-17/Domain: signal sequence #status predicted <SIG>
F:23,36,43,56,75,382,454,455,468,473/Modified site: 3',4'-dihydroxyphenylalanine (Tyr) #

Query Match 55.4%; Score 41; DB 2; Length 473;
Best Local Similarity 41.7%; Pred. No. 22;
Matches 5; Conservative 5; Mismatches 2; Indels 0; Gaps 0;

QY 3 CDSSECKSSPRC 14
| : : : : :
Db 86 CKPNOCKNSRC 97

RESULT 10
S32659
integrin beta 2 chain (CD18) - chicken
N:Alternate names: CD18 protein
C:Species: Gallus gallus (chicken)
C>Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 08-Nov-1996
C:Accession: I50660; S32659
R:Bilsland, C.A.; Springer, T.A.
J. Leukoc. Biol. 55, 501-506, 1994
A:Title: Cloning and expression of the chicken CD18 cDNA.
A:Reference number: I50660; MUID:94194252
A:Accession: I50660
A>Status: Preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: mRNA
A:Residues: 1-772 <BI2>
A:Cross-references: EMBL:X71786; NID:g297566; PID:g297567
C:Superfamily: integrin beta chain

Query Match 56.8%; Score 42; DB 2; Length 772;
Best Local Similarity 50.0%; Pred. No. 22;
Matches 6; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 HCDSECKSSPR 13
| : : : : :
Db 557 RCDGCECKCTPK 568

RESULT 11
SMKD25
metallothionein 2 - mud crab
C:Species: Scylla serrata (mud crab)
C>Date: 19-Feb-1984 #sequence_revision 19-Feb-1984 #text_change 13-Sep-1996
C:Accession: A03284
R:Lerch, K.; Ammer, D.; Olafson, R.W.
J. Biol. Chem. 257, 2420-2426, 1982
A:Title: Crab metallothionein. Primary structures of metallothioneins 1 and 2.
A:Reference number: A92363; MUID:82142340
A:Accession: A03284
A:Molecule type: protein

A:Residues: 1-57 <LER>
C:Superfamily: metallothionein
C:Keywords: metal binding

Query Match 48.6%; Score 36; DB 1; Length 57;
Best Local Similarity 41.7%; Pred. No. 26;
Matches 5; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 3 CDSSECKSSPRC 14
| : : : : :
Db 11 CREGECKTGCK 22

RESULT 12
A35844
Xotch protein - African clawed frog
C:Species: Xenopus laevis (African clawed frog)
C>Date: 12-Oct-1990 #sequence_revision 12-Oct-1990 #text_change 14-Aug-1998
C:Accession: A35844
R:Coffman, C.; Harris, W.; Kintner, C.
Science 249, 1438-1441, 1990
A:Title: Xotch, the Xenopus homolog of Drosophila notch.
A:Reference number: A35844; MUID:90385285
A:Accession: A35844
A>Status: Preliminary; nucleic acid sequence not shown; not compared with conceptual
A:Molecule type: mRNA
A:Residues: 1-2524 <COF>
C:Superfamily: unassigned ankyrin repeat proteins; ankyrin repeat homology; EGF homol
C:Keywords: transmembrane protein
F:222-254/Domain: EGF homology <EGF>
F:1924-1956/Domain: ankyrin repeat homology <AN1>
F:1957-1989/Domain: ankyrin repeat homology <AN2>
F:1991-2023/Domain: ankyrin repeat homology <AN3>
F:2024-2056/Domain: ankyrin repeat homology <AN4>
F:2057-2089/Domain: ankyrin repeat homology <AN5>

Query Match 59.5%; Score 44; DB 2; Length 2524;
Best Local Similarity 53.8%; Pred. No. 27;
Matches 7; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 3 CDSSECKSSPRCK 15
| : : : : :
Db 833 CAGSPCKNGGRCK 845

RESULT 13
A46019
gene Notch-1 protein - mouse
C:Species: Mus musculus (house mouse)
C>Date: 22-Sep-1993 #sequence_revision 18-Nov-1994 #text_change 14-Aug-1998
C:Accession: A46019
R:del Amo, F.F.; Gendron-Maguire, M.; Swiatek, P.J.; Jenkins, N.A.; Copeland, N.G.; G
Genomics 15, 259-264, 1993
A:Title: Cloning, analysis, and chromosomal localization of Notch-1, a mouse homolog
A:Reference number: A46019; MUID:93194170
A:Accession: A46019
A>Status: Preliminary; not compared with conceptual translation
A:Molecule type: nucleic acid
A:Residues: 1-2531
A:Cross-references: GB:Z11886; GB:S47228; NID:g288502; PID:g288503
A>Note: sequence extracted from NCBI backbone (NCBIP:127318)
C:Superfamily: unassigned ankyrin repeat proteins; ankyrin repeat homology; EGF homol
F:757-788/Domain: EGF homology <EGF>
F:1917-1948/Domain: ankyrin repeat homology <AN1>
F:1949-1981/Domain: ankyrin repeat homology <AN2>
F:1983-2015/Domain: ankyrin repeat homology <AN3>
F:2016-2048/Domain: ankyrin repeat homology <AN4>
F:2049-2081/Domain: ankyrin repeat homology <AN5>

Query Match 59.5%; Score 44; DB 2; Length 2531;

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Best Local Similarity 58.3%; Pred. No. 27;
Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 3 CDSSECKSSPRC 14
   |||: |||: ||
Db 1063 CDSAPCKNGRC 1074

RESULT 14
S44787
D2007.1 protein - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 14-Sep-1994 #sequence_revision 12-May-1995 #text_change 09-Sep-1997
C:Accession: S44787
R:Favell, A.D.
submitted to the EMBL Data Library, May 1993
A:Description: Sequence of the C. elegans cosmid D2007.
A:Reference number: S44617
A:Accession: S44787
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-111 <FAV>
A:Cross-references: EMBL:L16560; NID:g289666; PID:g289668
C:Genetics:
A:Introns: 48/1; 84/3

Query Match 50.0%; Score 37; DB 2; Length 111;
Best Local Similarity 33.3%; Pred. No. 30;
Matches 5; Conservative 7; Mismatches 3; Indels 0; Gaps 0;

QY 1 QHCDSECKSSPRCK 15
   ::||:: |||: ||
Db 91 DQGNACCKTSEQCR 105

RESULT 15
S36032
thrombin inhibitor haemadin - terrestrial leech (Haemadipsa sylvestris)
C:Species: Haemadipsa sylvestris
C:Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 09-Sep-1997
C:Accession: S36032; B46057
R:Kroger, B.
submitted to the EMBL Data Library, January 1993
A:Reference number: S36032
A:Accession: S36032
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-77 <KRV>
A:Cross-references: EMBL:Z19864; NID:g298108; PID:g298109
R:Strube, K.H.; Kroger, B.; Bialojan, S.; Otte, M.; Dödt, J.
J. Biol. Chem. 268, 8590-8595, 1993
A:Title: Isolation, sequence analysis, and cloning of haemadin. An anticoagulant peptide
A:Reference number: A46057; MUID:93232009
A:Accession: B46057
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-77 <STR>
A>Note: sequence extracted from NCBI backbone (NCBIN:129612, NCBI:P:129614)

Query Match 48.6%; Score 35; DB 2; Length 77;
Best Local Similarity 30.8%; Pred. No. 33;
Matches 4; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

QY 1 QHCDSECKSSPR 13
   |||: |||: ||
Db 50 QSCNDGQCGDPK 62

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: May 4, 1999, 12:32:37 ; Search time 27.23 Seconds
(without alignments)
11.141 Million cell updates/sec

Title: US-09-037-460-2_COPY_30_44

Perfect score: 74
Sequence: 1 QHCDSECKSSPRCK 15

Scoring table: PAM150

Searched: 162890 seqs, 20225328 residues

Database: A_Geneseq_34:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	74	100.0	184	1 R98994	Vascular TBP-like
2	39.5	53.4	50	1 R96122	Leech derived fahs
3	39.5	53.4	50	1 R96123	Leech derived fahs
4	42	56.8	139	1 R95596	Factor X light cha
5	44	59.5	467	1 W40283	Human Factor X pro
6	42	56.8	250	1 R13675	Factor X-LACI hydr
7	43	58.1	400	1 R32501	Beta-adrenergic re
8	37	50.0	52	1 W80576	Human factor VII E
9	37	50.0	53	1 R40210	Sequence of yeast
10	36	48.6	39	1 R37437	Thrombin inhibitor
11	37	50.0	61	1 P94625	CUP1 translation p
12	37	50.0	61	1 P50776	Sequence encoded b
13	37	50.0	61	1 P30006	Sequence of yeast
14	37	50.0	61	1 R54087	Metallothionein. P
15	42	56.8	436	1 R22513	Truncated precursor
16	42	56.8	437	1 R8468	Human two chain fa
17	42	56.8	437	1 R37402	Factor X. Factor X
18	42	56.8	448	1 R35762	Factor X (X). Seri
19	42	56.8	448	1 W66092	Human factor X var
20	34	45.9	21	1 W00600	Heparan sulphate p
21	44	59.5	1055	1 W44298	Human serrate 2 pr
22	42	56.8	488	1 R22512	Mutated precursor
23	42	56.8	488	1 R22511	Human Factor Xai
24	42	56.8	488	1 W76219	Human Factor X pro
25	42	56.8	488	1 W76216	Human Factor X pro
26	42	56.8	488	1 W76217	Human Factor X pro
27	42	56.8	488	1 W76218	Human Factor X pro
28	37	50.0	70	1 P94628	CUP1 translation p
29	37	50.0	70	1 P94627	CUP1 translation p
30	44	59.5	1212	1 W44299	Human serrate 2. H
31	44	59.5	1257	1 W05834	Human serrate-2 (H
32	37	50.0	85	1 R62821	Deduced sequence o
33	37	50.0	86	1 R62820	Recombinant factor
34	41	55.4	473	1 R86869	Adhesive protein.
35	37	50.0	110	1 R62622	Deduced sequence e
36	36	48.6	77	1 R37438	Thrombin inhibitor
37	37	50.0	148	1 W68344	Protein encoded by
38	37	50.0	133	1 R62516	Deduced sequence e
39	37	50.0	153	1 R62624	Deduced sequence o
40	39	52.7	388	1 R25698	Murine adrenergic
41	37	50.0	182	1 R62619	Deduced sequence o
42	30	40.5	12	1 R64297	Factor-VII-derived
43	30	40.5	12	1 R67863	FVII/TF multi-prot

44 39 52.7 400 1 R54992 Murine beta-3 adre
45 33 44.6 44 1 R96225 Novel growth facto

ALIGNMENTS

RESULT 1
R98994
ID R98994 standard; Protein; 184 AA.
AC R98994; 1996 (first entry)
DE Vascular IBP-like growth factor.
KW Vascular IBF-like growth factor; VIGF;
KW Insulin-like growth factor binding protein; agonist; antagonist;
KW muscle wastage; osteoporosis; implant fixation; wound healing;
KW therapy; diagnosis.
OS Homo sapiens.
FH Key Location/Qualifiers
FT peptide 1..21
FT /label= Sig_peptide
PN WO9617931-A1.
PD 13-JUN-1996.
PF 09-DEC-1994; U14388.
PR 09-DEC-1994; WO-U14388.
PI (HUMA-) HUMAN GENOME SCI INC.
PI Hastings GA, Rosen CA;
DR WPI: 96-287176/29.
DR N-PSDB; T34991.
PT Human vascular insulin-like growth factor binding protein-like
PT growth factor, and its nucleic acid sequence and (antagonists -
PT used, e.g. to treat muscle wasting diseases or aid implant fixation,
PT or limit excess connective tissue prodn. during wound healing.
PS Clam 4; Page 43-44; 61pp; English.
CC Human vascular insulin-like growth factor binding protein-like
CC growth factor (R98994), or VIGF, is a protein of primarily
CC vascular origin that is structurally related to the IBP and CCN
CC protein families. It can be expressed in e.g. E. coli, CHO or
CC (T34991), or its derivative, obtd. from human umbilical
CC endothelial cells. It is useful therapeutically e.g. for
CC treating muscle wasting diseases or osteoporosis, or can be used
CC to detect diseases associated with under- or over-expression of VIGF,
CC or to screen for antagonists useful during wound healing.
SQ Sequence 184 AA;

Query Match 100.0%; Score 74; DB 1; Length 184;
Best Local Similarity 100.0%; Pred. No. 0.00048;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 QHCDSECKSSPRCK 15
| | | | | | | | | | | | | | | | | |
Db 30 QHCDSECKSSPRCK 44

RESULT 2
R96122
ID R96122 standard; Peptide; 50 AA.
AC R96122;
DE 17-DEC-1996 (first entry)
DE Leech derived fahsin based protease inhibitor #2.
KW Protease inhibitor; Isoform; elastase; chymotrypsin; trypsin; leech;
KW tissue; secretion; saliva; fahsin; antidiabetic; diabetes mellitus;
KW blood clotting disorder; neutrophil function; emphysema;
KW rheumatoid arthritis; HIV infection; human immunodeficiency virus.
OS Limnatis nilotica.
PN WO9613585-A1.
PD 09-MAY-1996.
PF 27-OCT-1995; E04223.
PR 28-OCT-1994; EP-117053.
PR 14-MAR-1995; EP-103637.
PA (CLOD-) CLODICA SA.

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PI Voerman G;
DR WPI; 96-239498/24.
PT New protease inhibitors from the leech Limnatis nilotica - for
PT treating, e.g. blood clotting disorders, HIV infection, diabetes
PT mellitus etc.
PS Claim 3; Page 26; 41pp; English.
CC The protease inhibitor peptide isoforms given in R96121-23 are
CC elastase/chymotrypsin- and trypsin inhibitors which may be isolated
CC from leech tissue or leech secretions, e.g. saliva. These peptides
CC belong to the family of leech derived substances named fahsin's which
CC also have an antibiotic effect. The fahsin family of proteins comprise
CC 50/51 amino acids and occur in various isoforms. These peptides are
CC useful in the treatment of diabetes mellitus, blood clotting disorders,
CC disorders of neutrophil function, e.g. emphysema, rheumatoid arthritis,
CC HIV infection and other immunological and inflammatory diseases.
SQ Sequence 50 AA;

Query Match      53.4%; Score 39.5; DB 1; Length 50;
Best Local Similarity 43.8%; Pred. No. 11;
Matches 7; Conservative 4; Mismatches 4; Indels 1; Gaps 1;

QY 1 QHCDSECKSSP-RCK 15
Db 13 QLCVDGQCKCTPIRCR 28

RESULT 3
R96123
ID R96123 standard; Peptide; 50 AA.
AC R96123;
DE Leech derived fahsin based protease inhibitor #3.
KW Protease inhibitor; Isoform; elastase; chymotrypsin; trypsin; leech;
KW tissue; secretion; saliva; fahsin; antibiotic; diabetes mellitus;
KW blood clotting disorder; neutrophil function; emphysema;
KW rheumatoid arthritis; HIV infection; human immunodeficiency virus.
OS Limnatis nilotica.
PN W09613585-A1.
PD 09-MAY-1996.
PF 27-OCT-1995; E04223.
PR 28-OCT-1994; EP-117053.
PR 14-MAR-1995; EP-103637.
PA (CLOD-) CLODICA SA.
PI Voerman G;
DR WPI; 96-239498/24.
PT New protease inhibitors from the leech Limnatis nilotica - for
PT treating, e.g. blood clotting disorders, HIV infection, diabetes
PT mellitus etc.
PS Claim 3; Page 26; 41pp; English.
CC The protease inhibitor peptide isoforms given in R96121-23 are
CC elastase/chymotrypsin- and trypsin inhibitors which may be isolated
CC from leech tissue or leech secretions, e.g. saliva. These peptides
CC belong to the family of leech derived substances named fahsin's which
CC also have an antibiotic effect. The fahsin family of proteins comprise
CC 50/51 amino acids and occur in various isoforms. These peptides are
CC useful in the treatment of diabetes mellitus, blood clotting disorders,
CC disorders of neutrophil function, e.g. emphysema, rheumatoid arthritis,
CC HIV infection and other immunological and inflammatory diseases.
SQ Sequence 50 AA;

Query Match      53.4%; Score 39.5; DB 1; Length 50;
Best Local Similarity 43.8%; Pred. No. 11;
Matches 7; Conservative 4; Mismatches 4; Indels 1; Gaps 1;

QY 1 QHCDSECKSSP-RCK 15
Db 13 QLCVDGQCKCTPIRCR 28

RESULT 4
R95596
ID R95596 standard; protein; 139 AA.
AC R95596;
DE 17-DEC-1996 (first entry)
KW Factor X; light chain.
KW Factor X; light chain; human; heavy chain; Factor Xa; prothrombinase;
KW activating enzyme; blood factor; immunoaffinity chromatography; therapy;
KW antigen; inhibitor; activated resin; coagulation disorder; factor II;
KW vasculature function disorder; factor V; factor IX; factor XI; protein C;
KW factor XII; factor VII; protein S; fibrinogen.
OS Homo sapiens.
PH Key Location/Qualifiers
FT disulfide_bond 17..22
FT domain 17..22
FT /note= "GLA domain"
FT disulfide_bond 50..61
FT domain 50..124
FT /note= "growth factor domains"
FT disulfide_bond 55..70
FT misc_difference 63
FT /note= "unspecified amino acid, represented in
FT specification as beta"
FT disulfide_bond 72..81
FT disulfide_bond 89..100
FT disulfide_bond 96..109
FT disulfide_bond 111..124
FT disulfide_bond 132
FT /note= "disulfide bond to residue 160 of Factor X heavy
FT chain (see W05820), or to residue 108 of Factor
FT Xa heavy chain (see R95597 and R95598)."
PN W09613274-A1.
PD 09-MAY-1996.
PF 27-OCT-1995; U13940.
PR 28-OCT-1994; US-330978.
PA (CORP-) COR THERAPEUTICS INC.
PI King R8;
DR WPI; 96-239270/24.
PT Prepn. of an inhibited form of an activated blood factor, e.g.
PT factor X - by treating partially purified blood factor preps. with
PT an activating factor and an inhibiting factor
PS Disclosure; Fig 1; 45pp; English.
CC This sequence represents the light chain of human Factor X (the heavy
CC chain is represented by W05820). Factor X must be activated to Factor Xa
CC before the protease is incorporated into the prothrombinase complex. In
CC Factor Xa the light chain sequence is identical to the Factor X light
CC chain, and the heavy chain is a truncated version of the Factor X heavy
CC chain. An inhibited form of activated Factor X is prepared by the method
CC of the invention. In this method, a partially purified preparation
CC containing the blood factor is treated to convert the factor into an
CC activated form (using an immobilised activating enzyme). The activated
CC form is then converted into an inhibited form in a single step, and the
CC inhibited factor is recovered. The inhibited blood factor is recovered
CC by immunoaffinity chromatography using an antigen specific monoclonal
CC antibody coupled to an activated resin (such as agarose), or an anion
CC exchange column with an anion-exchange group linked to a naturally
CC derived polysaccharide or a synthetically derived polymeric matrix. The
CC activated resin used preferably uses activation chemistry selected from
CC tressyl, azactone, aldehyde, hydrazide, N-hydroxy succinimide or triazine.
CC This method produces a highly purified preparation of an inhibited form
CC (either permanently or transiently inhibited) of an activated blood
CC factor in high yield. The factors produced can be used in the treatment
CC of coagulation disorders, or disorders of vasculature function.
SQ Sequence 139 AA;

Query Match      56.8%; Score 42; DB 1; Length 139;
Best Local Similarity 33.3%; Pred. No. 12;
Matches 5; Conservative 6; Mismatches 4; Indels 0; Gaps 0;

QY 1 QHCDSECKSSPRCK 15
Db 48 DQCTSPCQNGRCK 62

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FT	peptide	/label= Xlc	
FT		1. .40	
FT	domain	/label= prepro_leader	
FT		55. 64	
FT	domain	/label= GLA_domain	
FT		89. .150	
FT	domain	/label= growth_factor_domains	
FT		172. .250	
FT	domain	/label= kunitz_domain	
FT	disulfide_bond	57. .62	
FT	disulfide_bond	90. .101	
FT	disulfide_bond	95. .110	
FT	disulfide_bond	112. .121	
FT	disulfide_bond	129. .140	
FT	disulfide_bond	136. .149	
FT	disulfide_bond	151. .164	
FT	disulfide_bond	186. .236	
FT	disulfide_bond	195. .219	
FT	disulfide_bond	211. .232	
PN	EP-439442-A.		
PD	31-JUL-1991.		
PF	21-JAN-1991; 870008.		
PR	25-JAN-1990; US-470289.		
PP	(UNIW) UNIV OF WASHINGTON.		
PA	Girard TJ, Broze GJ;		
PI	WPI; 91-224839/31.		
DR	WPI; 91-224839/31.		
DR	PFSDB; Q12776.		
PT	New factor X-LACI hybrid protein - comprises light chain of		
PT	factor X and LACI's first kunitz domain for use as anticoagulant		
PS	disclosure; Page 12-14; 17pp; English.		
CC	The protein is used as a blood coagulation inhibitor in mammals. It		
CC	is believed to mimic the Xa/LACI complex in binding to and		
CC	inhibiting VIIa/tissue factor. LACI inhibits via a novel feedback		
CC	mechanism requiring generation of Xa (a prod. of VIIa/TF activity);		
CC	XlcLACIKI inhibits VIIa/TF activity directly.		
CC	The DNA allows prodn. of XlcLACIKI by introduction of the gene into		
CC	cells suitable for expression, e.g. E. coli or CHO cells.		
SQ	Sequence 250 AA;		

Query Match 56.8%; Score 42; DB 1; Length 250;
Best Local Similarity 33.3%; Pred. No. 19;
Matches 5; Conservative 6; Mismatches 4; Indels 0; Gaps 0;

RESULT 7
R32501
ID R32501 standard; protein; 400 AA.

DT	09-JUN-1993	(first entry)
DE	Beta-adrenergic receptor.	
DE	Fat cell specific; BAR; lipolysis; obesity; diagnosis;	
KW	thermogenesis; metabolism.	
OS	Rattus rattus.	
PN	US7783602-A.	
PD	15-DEC-1992.	
PF	11-NOV-1991; 783602.	
PP	01-NOV-1991; US-783602.	
PPA	(US\$) US DEPT HEALTH & HUMAN SERVICE.	
PI	Venter CJ;	
PI	WPI: 93-067426/08.	
DR	Fat cell specific beta- adrenergic receptor polypeptide	
PPT	for diagnosis of obesity due to inactive lipolysis	
PPT	Disclosure; Page 16; 20pp; English.	
PPS	A rat intracapsular brown adipose tissue cDNA library	
CCC	probes with DNA probes encoding human beta-1 and rat b	
CCC	adrenergic receptors under low stringency conditions.	
CCC	clones were found to be different from the rat and human	
CCC	and contained a single open reading frame of 1200 bp	
CCC		

CC protein shown, of 400 amino acids and mol. wt. 43 kD. The protein is
 CC the fat specific beta-adrenergic receptor and may be used in work on
 CC the thermogenesis process. Isolation of the gene for BAR allows the
 CC diagnosis and treatment of obesity and the testing of medications
 CC for their effectiveness in stimulating the thermogenesis metabolic
 CC response in obesity patients.
 SQ Sequence 400 AA;

Query Match 58.1%; Score 43; DB 1; Length 400;
 Best Local Similarity 58.3%; Pred. No. 21;
 Matches 7; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Oy 4 DSSECKSPRCK 15
 Db 182 EAQCHSNPRCK 193

RESULT 8
 W80576 ID W80576 standard; Protein; 52 AA.
 AC W80576; 16-DEC-1998 (first entry)
 DE Human factor VII EGF domain primary sequence.
 KW O-fucosyltransferase; epidermal growth factor; EGF; glycosylation;
 KW O-fucose; inhibitor; sensory neuron; retinal neuron; human; heart.
 OS Synthetic.
 OS Homo sapiens.

FH Key Location/Qualifiers
 FT Peptide 44..49 /note="histidine tag"
 FT W09833924-A1.
 PN 06-AUG-1998.
 PD 17-DEC-1997; U23401.
 PF 26-NOV-1997; US-978741.
 PR 31-JAN-1997; US-792498.
 PA (GETH) GENENTECH INC.
 PI Spellman MW, Wang Y;
 DR WPI: 98-437477/37.
 PT Human O-fucosyltransferase able to glycosylate epidermal growth
 PT factor domains - useful for diagnosis and treatment of diseases
 PT involving overexpression of the enzyme
 PS Example 2; Page 39; 90pp; English.
 CC This represents the primary sequence of the first EGF domain from human
 CC factor VII contained in a plasmid construct. The invention provides a
 CC human heart O-fucosyltransferase enzyme that can glycosylate an epidermal
 CC growth factor (EGF) domain of a polypeptide with an activated O-fucose
 CC residue. Inhibitors of O-fucosyltransferase, e.g. mutants with increased
 CC affinity for the EGF domains, are used in diagnosis and treatment of
 CC conditions associated with overexpression of O-fucosyltransferase, to
 CC promote survival of sensory (retinal) neurons. Probes based on EGF domain
 CC polypeptide are used to detect gene amplification and expression. The
 CC expression can also be determined at the protein level using antibodies
 CC specific for O-fucosyltransferase.
 SQ Sequence 52 AA;

Query Match 50.0%; Score 37; DB 1; Length 52;
 Best Local Similarity 33.3%; Pred. No. 26;
 Matches 5; Conservative 6; Mismatches 4; Indels 0; Gaps 0;

Oy 1 OHCDSECKSPRCK 15
 Db 5 DQCESNPLNGSGCK 19

RESULT 9
 R40210 ID R40210 standard; protein; 53 AA.
 AC R40210.
 DT 04-FEB-1994 (first entry)
 DE Sequence of yeast class II metallothionine.
 KW Metallothionine; yeast; class II.

OS Saccharomyces cervisiae.
 PN DE4212134-A.
 PD 19-AUG-1993.
 PF 10-APR-1992; 212134.
 PR 17-FEB-1992; GB-003299.
 PA (INDE-) INDENA SPA.
 PI Bombardelli E, Ponzone C, Puglisi PP;
 DR WPI: 93-265710/34.
 PT Topical compsn. for protecting tissue e.g. skin - against toxic
 PT heavy metals, contg. metal-complexing protein with high cysteine
 PT content
 PS Disclosure; Page 3; 7pp; German.
 CC Class II metallothionines have the same potential to form
 CC thio-metal complexes and clusters as Class II metallothionines.
 CC This class comprises metallothionines whose primary structure
 CC bears only a slight, or no, resemblance to those of mammals.
 SQ Sequence 53 AA;

Query Match 50.0%; Score 37; DB 1; Length 53;
 Best Local Similarity 30.8%; Pred. No. 27;
 Matches 4; Conservative 6; Mismatches 3; Indels 0; Gaps 0;

Oy 2 HCDSECKSPRCK 14
 Db 8 QCQCGSKNEQC 20

RESULT 10
 R37437 ID R37437 standard; Protein; 39 AA.
 AC R37437;
 DE 30-SEP-1993 (first entry)
 DE Thrombin inhibitor C-terminal (AS 19-57).
 KW Primer; PCR; thrombosis; arterial reocclusion; blood; thrombin;
 KW hirudin.
 OS Haemadipsa sylvestris.
 PN DE4209110-A.
 PD 27-MAY-1993.
 PF 20-MAR-1992; 209110.
 PR 26-NOV-1991; DE-138698.
 PR 20-MAR-1992; DE-209110.
 PA (BADI) BASF AG.
 PI Bialojan S, Friedrich T, Kroeger B, Strube K;
 DR WPI: 93-176720/22.
 DR N-PSDB: Q42433.
 PT New proteins obtd. from Haemadipsa Sylvestris - used as thrombin
 PT inhibitors for treatment and prevention of thrombosis and
 PT arterial re-occlusion
 PS Example 6; Page 8; 14pp; German.
 CC Example 6 describes the cloning of DNA encoding the thrombin
 CC inhibiting protein. A cDNA library is produced from H. sylvestris
 CC DNA. Primers (i - Q42426) and (ii - Q42427) were based on peptide
 CC fragments corresp. to amino acids 4-11 and 19-28 respectively.
 CC The 3' primers were A-B-T18 (Q42428), A (derived from A-B-T18 -
 CC Q42429) and B (derived from A-B-T18 - Q42430). A first sequence
 CC was obtained (Q42433). A further PCR cycle was performed using
 CC primers (iii - Q42431) and (iv - Q42432). The sequence
 CC given in Q42434 was obtained.
 SQ Sequence 39 AA;

Query Match 48.6%; Score 36; DB 1; Length 39;
 Best Local Similarity 30.8%; Pred. No. 29;
 Matches 4; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

Oy 1 OHCDSECKSPRCK 13
 Db 12 QSCNDGQCSGDPK 24

RESULT 11
 P94625

ID P94625 standard; protein; 61 AA.
 AC P94625;
 DT 22-JUN-1990 (first entry)
 DE CUP1 translation product of plasmid M13mp18/CUP1.
 KW Copper resistance gene; expression system; yeast; cassette; ds.
 OS Saccharomyces cerevisiae
 PN AU8815845-A.
 PD 10-NOV-1988.
 PF 7-MAY-1987; 15845.
 PR 7-MAY-1987; AU-015845.
 PR 6-MAY-1988; AU-001788.
 PA (CSIR) Commonwealth Scient Org.
 PI Macreadie IG, Vaughan PR, Azad AA;
 DR WPI; 89-069074/10.
 DR N-PSDB; N90634.
 PT Gene expression system -
 PT contains yeast copper resistance gene and multiple cloning site
 PT and is movable as a cassette.
 PS Disclosure; P; English.
 CC Expression vector carrying the yeast copper resistance genes allows for
 CC the expression of foreign proteins with a copper induced promoter and in
 CC the absence of their own start codon.
 SQ Sequence 61 AA;

Query Match 50.0%; Score 37; DB 1; Length 61;
 Best Local Similarity 30.8%; Pred. No. 30;
 Matches 4; Conservative 6; Mismatches 3; Indels 0; Gaps 0;
 QY 2 HCDSSSECKSSPRC 14
 DB 16 QCQCGSKNNEQC 28

RESULT 12
 ID P50776
 AC P50776 standard; Protein; 61 AA.
 DT 30-NOV-1991 (first entry)
 DE Sequence encoded by yeast copper metallothionein gene in
 DE YEP36 insert.
 KW Metal poisoning therapy; chelating agent;
 KW industrial waste treatment.
 OS Saccharomyces cerevisiae.
 PN EP-134773-A.
 PD 20-MAR-1985.
 PF 07-AUG-1984; 870112.
 PR 08-AUG-1983; US-520668.
 PR 02-MAR-1984; US-584657.
 PA (SMIK) SMITHKLINE BECKMAN CORP.
 PI Butt TR;
 DR WPI; 85-070084/12.
 DR N-PSDB; N50529.
 PT Gene coding for yeast copper metallothionein - isolated from
 PT yeast by hybridisation and useful for micro-organism
 PT transformation
 PS Disclosure; Page 4-6; 25pp; English.
 CC The gene is present in a Sau 3A-Sau 3A fragment of chromosomal DNA
 CC in a Cu-resistant (imm CuSO4) mutant. The promoter region is present
 CC in a Sau 3A-Kpn I, XbaI-Kpn I or Rsa I-Kpn I fragment. Recombinant
 CC DNA molecules contg. the structural gene include YEP1, YEP2, YEP29,
 CC YEP36 and TRP20. (all claimed).
 SQ sequence 61 AA;

Query Match 50.0%; Score 37; DB 1; Length 61;
 Best Local Similarity 30.8%; Pred. No. 30;
 Matches 4; Conservative 6; Mismatches 3; Indels 0; Gaps 0;
 QY 2 HCDSSSECKSSPRC 14
 DB 16 QCQCGSKNNEQC 28

RESULT 13
 ID P30006
 AC P30006 standard; peptide; 61 AA.
 DT 25-APR-1992 (first entry)
 DE Sequence of yeast copper chelating metallothionein
 DE Cu-chelatin).
 KW Copper chelating metallothionein; heavy metal; copper-resistant;
 KW yeast vector; industrial waste.
 OS Saccharomyces cerevisiae strain B231-1-7Ba.
 PN EP-36491-A.
 PD 21-DEC-1983.
 PF 03-JUN-1983; 098143.
 PR 03-JUN-1982; US-384821.
 PA (REGC) UNIV OF CALIFORNIA.
 PI Fogel S, Welch JW, Karin M;
 DR WPI; 83-846233/51.
 DR N-PSDB; N30011.
 PT Pure yeast copper chelatin - prodn. by recombinant DNA methods
 PT from DNA sequences
 PS Disclosure; Page 7; 16pp; English.
 CC The inventors claim yeast copper chelatin, its fragments and
 CC analogues and DNA encoding them. Cu-chelatin is useful as a
 CC chelating agent for removal of heavy metals, esp. copper, from
 CC industrial wastes, etc. Plasmids isolated from hosts transformed to
 CC copper resistant are useful in prepn. of hybridisation probes. A
 CC yeast expression system contg. the Cu-chelatin DNA and upstream
 CC regulatory sequence together with a foreign gene and regulatory
 CC signals is also claimed.
 SQ Sequence 61 AA;

Query Match 50.0%; Score 37; DB 1; Length 61;
 Best Local Similarity 30.8%; Pred. No. 30;
 Matches 4; Conservative 6; Mismatches 3; Indels 0; Gaps 0;
 QY 2 HCDSSSECKSSPRC 14
 DB 16 QCQCGSKNNEQC 28

RESULT 14
 ID R54087
 AC R54087 standard; Protein; 61 AA.
 DT 01-FEB-1995 (first entry)
 DE Metallothionein.
 KW Metallothionein; CUP1; promoter; yeast; copper; expression;
 OS Synthetic.
 PN EP-603128-A.
 PD 22-JUN-1994.
 PF 07-DEC-1993; 810853.
 PR 15-DEC-1992; EP-811005.
 PA (CIBA) CIBA GEIGY AG.
 PA (UCPG-) UCP GEN-PHARMA AG.
 PI Fuerst P, Heim J, Hottiger T, Pohlrig G;
 DR WPI; 94-193627/24.
 DR N-PSDB; Q64145.
 PT Production of polypeptide(s) with improved stability and
 PT increased yields using engineered yeast cells transformed with
 PT plasmids carrying a CUP1 gene, and adding copper to the culture
 PT medium.
 PS Disclosure; Page 20-21; 36pp; English.
 CC Example 1 describes the construction of plasmid pHE112R, a 2 micron
 CC plasmid contg. the GAPFLP-hirudin expression cassette and the full
 CC CUP1 gene. A 1.3 kb BamHI fragment contg. the full metallothionein
 CC encoding gene - CUP1, is isolated from plasmid YEP3362xSst (Wright,
 CC C.F. et al., Nucleic Acids Res. 14 (1986), 8489-8499). YEP3362xSst
 CC is digested with BamHI, the 1.3 kb fragment isolated, purified and
 CC ligated with BamHI cut pUC19. The resulting plasmid is named pHE105.
 CC The 1.3 kb fragment (see Q64145) is ligated into SmaBI-cut

CC PDP34/GAPFL-YHIR. E. coli is transformed with the resulting
CC plasmid pHEL12R.
SQ Sequence 61 AA;

Query Match 50.0%; Score 37; DB 1; Length 61;
Best Local Similarity 30.8%; Pred. No. 30;
Matches 4; Conservative 6; Mismatches 3; Indels 0; Gaps 0;
QY 2 HCDSECKSSPRC 14
Db 16 QCCGCKRNEQC 28
:|:|:|:|:|

RESULT 15
R22513
ID R22513 standard; Protein; 436 AA.
AC R22513;
DT 28-JUL-1992 (first entry)
DE Truncated precursor of human Factor Xa.
KW Mutant; prothrombinase complex; proteolytic; precursor; thrombosis;
KW inflammation; restenosis; transplantation; haemophilia; antibodies.
OS Homo sapiens.
PN WO9204378-A.
PD 19-MAR-1992.
PF 04-SEP-1991; U06337.
PR 04-SEP-1990; US-578646.
PA (COR-) COR THERAPEUTICS IN.
PI Wolf D;
DR WPI; 92-114303/14.
PT New analogues of Factor Xa peptide - useful for treating
PT haemophilia, thrombosis, inflammation and transplant
PT complications, for in-vivo diagnosis
PS Claim 7; Fig 1; 59pp; English.
CC The full length cDNA of human factor X (Mpl9X) was converted to
CC encode a truncated form of human Factor X, designated 'rx', by deletion
CC of the activation peptide by oligonucleotide site directed mutagenesis.
CC An oligonucleotide was used to align Arg 142 following the
CC C-terminus of the Factor X light chain with ile 53 of the Factor X
CC activation peptide (1st residue of the heavy chain). When expressed
CC in CHO cells the truncated peptide was cleaved endogenously.
CC Modified Factor Xa was further produced by acylation e.g. with the
CC p-nitrophenyl ester of p-toluoylic acid. Factor 'rx' is used to
CC treat or prevent thrombosis; inflammation; restenosis or complications
CC of transplantation. It is also used in treatment of adult respiratory
CC distress syndrome and haemophilia. The modified Factor X has no
CC proteolytic activity and interferes with the ability of endogenous
CC factor Xa to convert prothrombin to thrombin. Antibodies reactive
CC with Factor 'rx' are passive therapeutic agents and used for diagnosis.
CC See also R22512.
SQ Sequence 436 AA;

Query Match 56.8%; Score 42; DB 1; Length 436;
Best Local Similarity 33.3%; Pred. No. 31;
Matches 5; Conservative 6; Mismatches 4; Indels 0; Gaps 0;
QY 1 QHCDSECKSSPRCK 15
Db 88 DQCETSPQONQCK 102
:|:|:|:|:|

Search completed: May 4, 1999, 12:32:39
Job time: 9786 sec

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OM protein - protein search, using sw model

Run on: May 4, 1999, 06:22:08 ; Search time 25.07 Seconds
(without alignments)
243.559 Million cell updates/sec

Title: US-09-037-460-2_COPY_1_163
Perfect score: 820
Sequence: 1 MKSVLLTLLVPAHLVA...NRFSVLTREHDWASGDNIVR 163

Scoring table: PAM150

Searched: 116738 seqs, 37460341 residues

Database : PIR_58:*
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	141.5	17.3	351	2 S20078	NOV protein - chic
2	123.5	15.1	357	2 I38069	gene novH protein
3	115.5	14.1	258	2 A45403	insulin-like growth
4	114.5	14.0	271	2 I48604	insulin-like growth
5	112	13.7	348	2 A40578	beta IG-M2 protein
6	112	13.7	348	2 A53228	fisp-12 protein pr
7	110.5	13.5	271	2 JCI463	insulin-like growth
8	111	13.5	375	2 A41428	insulin-like growth
9	109.5	13.4	272	2 A53748	CEP-10 protein pre
10	109	13.3	349	2 A40551	insulin-like growth
11	107	13.0	237	2 I47031	connective tissue
12	106.5	13.0	258	2 G01662	insulin-like growth
13	106.5	13.0	258	2 B37252	insulin-like growth
14	104.5	12.7	266	2 A35037	insulin-like growth
15	104.5	12.7	271	2 JCI4584	insulin-like growth
16	103.5	12.6	254	2 I48599	insulin-like growth
17	103.5	12.6	254	2 JCI484	insulin-like growth
18	103.5	12.6	379	2 A35669	gene CYR61 protein
19	101.5	12.4	291	1 IOH03	insulin-like growth
20	101.5	12.4	291	1 JN0084	insulin-like growth
21	100.5	12.3	291	2 I48602	insulin-like growth
22	98.5	12.0	292	2 A36748	insulin-like growth
23	98	12.0	251	2 A55035	insulin-like growth
24	103	12.6	1801	1 MWR5	cysteine-rich prot
25	101.5	12.4	1138	1 S24066	laminin beta-2 cha
26	97	11.8	310	2 A20967	protein-tyrosine k
27	97	11.8	317	2 A60967	insulin-like growth
28	94.5	11.5	259	1 IOH01	insulin-like growth
29	90.5	11.0	81	2 A45320	transglutaminase s
30	90.5	11.0	1136	1 S57845	protein-tyrosine k
31	94.5	11.5	1134	1 JN0711	insulin-like growth
32	87.5	10.7	111	2 B45403	insulin-like growth
33	94.5	11.5	1786	1 MWR5B1	laminin beta-1 cha
34	93.5	11.4	1816	1 S68960	laminin alpha-4 ch
35	92.5	11.3	1797	2 A56677	insulin-like growth
36	92	11.2	1786	1 MWR5B1	laminin beta-2 cha
37	86.5	10.5	328	2 A41927	laminin beta-1 cha
38	85.5	10.4	254	2 I48603	insulin-like growth
39	94	11.5	5147	1 IJFTFM	cadherin-related t

40 88 10.7 728 2 I50719
41 90 11.0 1639 1 MMFFB2
42 84.5 10.3 282 2 S50031
43 84.5 10.3 304 2 A33274
44 84.5 10.3 305 2 I48601
45 84.5 10.3 305 2 JN0508

ALIGNMENTS

RESULT 1
S20078
NOV protein - chicken
C:Species: Gallus gallus (chicken)
C:Date: 19-Feb-1994 #sequence_revision 10-Nov-1995 #text_change 10-Sep-1997
C:Accession: S20078
R:Joliot, V.; Martinierie, C.; Dambrine, G.; Plassiat, G.; Brisac, M.; Crochet, J.; P
Mol. Cell. Biol. 12, 10-21, 1992
A:Title: Proviral rearrangements and overexpression of a new cellular gene (nov) in m
A:Reference number: S20078; MUID:92107157
A:Accession: S20078
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-351 <JOL>
A:Cross-references: EMBL:X59284; NID:g63702; PID:g63703
C:Genetics:
A:Gene: NOV

Query Match 17.3%; Score 141.5; DB 2; Length 351;
Best Local Similarity 31.9%; Pred. No. 3.1e-08;
Matches 45; Conservative 17; Mismatches 38; Indels 41; Gaps 7;
QY 3 SVLLTLLVPAHLVAANNVAVDCPQCDSECKSSPRCK---RTVLDDGCGCVRCAA 59
Db 10 PVLLLLLLLLRCVEVSGR---EAAACPRCGRCAPGVPVAVLDGCGCLVCAR 65
QY 60 GRGETCYFTVSGMDGMKGGLRCOPSG-----EPFGEFGICK-----DCPYG-- 106
Db 66 QRGES-----CSPLPCDESGGLYCDRGPED---GGGAGICMVLEGNCVFDGM 111
QY 106 -----TFGMDRETNCOSG 120
Db 112 IYRNGETFPQCKYQCTCRDG 132

RESULT 2
I38069
gene novH protein - human
C:Species: Homo sapiens (man)
C:Date: 17-May-1996 #sequence_revision 17-May-1996 #text_change 28-Feb-1997
C:Accession: I38069
R:Martinierie, C.; Huff, V.; Joubert, I.; Badziach, M.; Saunders, G.; Strong, L.; Perb
Oncogene 9, 2729-2732, 1994
A:Title: Structural analysis of the human nov proto-oncogene and expression in Wilms
A:Reference number: I38069; MUID:94336229
A:Accession: I38069
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-357 <RES>
A:Cross-references: EMBL:X78351; NID:g587422; PID:g825696
C:Genetics:
A:Gene: novH
A:Introns: 28/3; 104/1; 188/1; 259/3

Query Match 15.1%; Score 123.5; DB 2; Length 357;
Best Local Similarity 31.8%; Pred. No. 2.8e-06;
Matches 42; Conservative 25; Mismatches 34; Indels 31; Gaps 9;
QY 7 LITLLVPAHLVA--AWSNNVAVDCPQCDSECKSSPRCK---RTVLDDGCGCVRCAA 61
Db 112 IYRNGETFPQCKYQCTCRDG 132

[illegible]

QY 6 LTTLLVLAHLVAWSNNVAVDCPOHDS--SECKSSPRCKRTVLD-DGCGCCRVCAARG 62
 Db 6 LVAALLAAGPGSLGDE-AHCPPCEKLARCPRPPVCGEELVREAGCGCCATCALGLG 64
 QY 63 ETCVTVSGMDGMKCGPLGRQPSNGD 90
 Db 65 MPC-----GVYTPRCGSLRCYPRGVE 87
 RESULT 13
 B37252
 insulin-like growth factor-binding protein 4 precursor - human
 N:Alternate names: IGFBP-4; inhibitory insulin-like growth factor-binding protein; in
 C:Species: Homo sapiens (man)
 C:Date: 28-Mar-1991 #sequence_revision 28-Mar-1991 #text_change 13-Sep-1998
 C:Accession: B37252; B39842; A36549; A60712; A54650; A49801; A34419
 R:Shimasaki, S.; Uchiyama, F.; Shimomura, M.; Ling, N.
 Mol. Endocrinol. 4, 1451-1458, 1990
 A:Title: Molecular cloning of the cDNAs encoding a novel insulin-like growth factor-b
 A:Reference number: A37252; MUID:91133415
 A:Accession: B37252
 A:Status: not compared with conceptual translation
 A:Molecule type: mRNA
 A:Residues: 1-258 <SHI>
 R:Kiefer, M.C.; Maslarz, F.R.; Bauer, D.M.; Zapf, J.
 J. Biol. Chem. 266, 9043-9049, 1991
 A:Title: Identification and molecular cloning of two new 30-kDa insulin-like growth f
 A:Reference number: A39842; MUID:91225006
 A:Accession: B39842
 A:Molecule type: mRNA
 A:Residues: 1-258 <KIE>
 A:Cross-references: GB:M62403; NID:g184815; PID:g184816
 R:Latour, D.; Mohan, S.; Linkhart, T.A.; Baylink, D.J.; Strong, D.D.
 Mol. Endocrinol. 4, 1806-1814, 1990
 A:Title: Inhibitory insulin-like growth factor-binding protein: cloning, complete seq
 A:Reference number: A36549; MUID:91186988
 A:Accession: A36549
 A:Molecule type: mRNA
 A:Residues: 1-50, 'A', 52-197, 'F', 199-258 <LAT>

R:Perkel, V.S.; Mohan, S.; Baylink, D.J.; Linkhart, T.A.
J. Clin. Endocrinol. Metab. 71, 533-535, 1990
A:Title: An inhibitory insulin-like growth factor binding protein (In-IGFBP) from human
A:Reference number: A60712
A:Accession: A60712
A:Molecule type: protein
A:Residues: 22-26, 'X', 28-29, 'X', 31-35 <PER>
R:Mohan, S.; Baylink, D.J.
Growth Regul. 1, 110-118, 1991
A:Title: Evidence that the inhibition of TE85 human bone cell proliferation by agents with
A:Reference number: A54650; MUID:93091814
A:Accession: A54650
A:Molecule type: protein
A:Residues: 22-29, 'X', 31-42 <MOH>
A:Experimental source: TE85 osteosarcoma cells
A:Note: sequence extracted from NCBI backbone (NCBIP:121076)
R:Culouscou, J.M.; Shoyab, M.
Cancer Res. 51, 2813-2819, 1991
A:Title: Purification of a colon cancer cell growth inhibitor and its identification as
A:Reference number: A49801
A:Accession: A49801
A:Molecule type: protein
A:Residues: 22-53 <CU>
A:Experimental source: colon adenocarcinoma cells
R:Mohan, S.; Bautista, C.M.; Wergedal, J.; Baylink, D.J.
Proc. Natl. Acad. Sci. U.S.A. 86, 8338-8342, 1989
A:Title: Isolation of an inhibitory insulin-like growth factor (IGF) binding protein from
A:Reference number: A34419; MUID:90046792
A:Accession: A34419
A:Molecule type: protein
A:Residues: 22-29, 'E', 31-32, 'A', 34-36 <MO>
C:Genetics:
A:Gene: GDB:IGFBP4
A:Cross-references: GDB:126811; OMIM:146733
A:Map position: 17q12-17q21
C:Superfamily: thyroglobulin type I repeat homology
C:Keywords: glycoprotein
F:1-21/Domain: signal sequence #status predicted <SIG>
F:22-258/Product: insulin-like growth factor-binding protein 4 #status experimental <MAI>
F:174-249/Domain: thyroglobulin type I repeat homology <THYI>
F:125/Binding site: carbohydrate (Asn) (covalent) #status experimental

Query Match 13.0%; Score 106.5; DB 2: Length 258;
Best Local Similarity 31.8%; Pred. No. 0.00015;
Matches 28; Conservative 22; Mismatches 29; Indels 9; Gaps 4;

Qy 6 LTTLLVPAHLVAANSNNYAVDCPOHCDSD--SECKSSPRCKRTVLD-DCGCCRVCAAGRG 62
Db 6 LVAALLAAGPGPSLGD-EAHCPCSEKILARCPVGCVELVREPGCGCCATCALGLG 64

Qy 63 ETCYFTVSGMDGKCGPLRCOPNSNGED 90
Db 65 MPC-----GVYTPRCGSLRCYPPRGVE 87

RESULT 14
A35037
Insulin-like growth factor-binding protein 3 homolog - pig
C:Species: Sus scrofa domestica (domestic pig)
C:Date: 20-Jul-1990 #sequence_revision 20-Jul-1990 #text_change 01-Dec-1995
C:Accession: A35037
R:Shimasaki, S.; Shimonaka, M.; Ui, M.; Inouye, S.; Shibata, F.; Ling, N.
J. Biol. Chem. 265, 2198-2202, 1990
A:Title: Structural characterization of a follicle-stimulating hormone action inhibitor
A:Reference number: A35037; MUID:90130475
A:Accession: A35037
A:Status: Preliminary
A:Molecule type: mRNA
A:Residues: 1-266 <SHI>
A:Cross-references: GB:J05228
C:Superfamily: insulin-like growth factor binding protein 1; thyroglobulin type I repeat
F:188-260/Domain: thyroglobulin type I repeat homology <THYI>

Query Match 12.7%; Score 104.5; DB 2: Length 266;
Best Local Similarity 31.3%; Pred. No. 0.00026;
Matches 26; Conservative 18; Mismatches 20; Indels 19; Gaps 4;

Qy 31 HCDSECK-----SSPRCKRTVLD-DCGCCRVCAAGRETCTRTVSGMDGKCGPG 80
Db 12 RCPCDARALAACAPPAAPCAELVREPGCGCCCLTALREGQAC-----GVYTERCGAG 66

Qy 81 LRCQPSNGE----DPFGEEFGIC 99
Db 67 LRCQPPGEPRLQALLDGRGIC 89

RESULT 15
JC4584
Insulin-like growth factor binding protein-5 - pig
C:Species: Sus scrofa domestica (domestic pig)
C:Date: 10-Apr-1996 #sequence_revision 24-May-1996 #text_change 10-Oct-1997
C:Accession: JC4584; G23734
R:White, M.E.; Diao, R.; Hathaway, M.R.; Mickelson, J.; Dayton, W.R.
Biochem. Biophys. Res. Commun. 218, 248-253, 1996
A:Title: Molecular cloning and sequence analysis of the porcine insulin-like growth f
A:Reference number: JC4584
A:Accession: JC4584
A:Molecule type: mRNA
A:Residues: 1-271 <WHI>
A:Cross-references: GB:U41340; MUID:g1173906; PID:g1173907
A:Experimental source: skeletal muscle
R:Shimasaki, S.; Gao, L.; Shimonaka, M.; Ling, N.
Mol. Endocrinol. 5, 938-948, 1991
A:Title: Isolation and molecular cloning of insulin-like growth factor-binding protei
A:Reference number: A23734; MUID:92049376
A:Accession: G23734
A:Molecule type: protein
A:Residues: 20-25, 'X', 27-28, 'X', 30-36, 'X', 38-39 <SHI>
C:Comment: This protein has essential roles in the regulation and coordination of ins
lays a role during myoblast proliferation and differentiation, and is important in th
C:Superfamily: thyroglobulin type I repeat homology
C:Keywords: differentiation; growth factor; skeletal muscle
F:1-19/Domain: signal sequence #status predicted <SIG>
F:20-271/Product: insulin-like growth factor binding protein-5 #status experimental <
F:191-262/Domain: thyroglobulin type I repeat homology <THYI>

Query Match 12.7%; Score 104.5; DB 2: Length 271;
Best Local Similarity 27.7%; Pred. No. 0.00026;
Matches 26; Conservative 20; Mismatches 27; Indels 21; Gaps 3;

Qy 4 VLLTTLVPAHLVAANSNNYAVDCPOHCDSDSECKSSPRCKRTVLD-----DCGCCRV 56
Db 7 LLLLAACAGPAQGLGSFV-----HCEPCDEKALSMCPSPGLGELVKDPCGCCMT 57

Qy 57 CAAGRETCTRTVSGMDGKCGPLRCOPNSNGED 90
Db 58 CALAEGOSC-----GVYTERCAOGLRCLPRQDEE 86

Search completed: May 4, 1999, 08:18:15
Job time: 6967 sec

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OM protein - protein search, using sw model

Run on: May 4, 1999, 09:49:33 ; Search time 27.23 Seconds
(without alignments)
121.070 Million cell updates/sec

Title: US-09-037-460-2_COPY_1_163
Perfect score: 820
Sequence: 1 MKSVLLTLLVPAHLVAW.....NRFVSLTEHDMASGDGNIVR 163

Scoring table: PAM150

Searched: 162890 seqs, 20225328 residues

Database : A_Geneseq_34.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	820	100.0	184	1 R98994	Vascular IBP-like
2	141.5	17.3	351	1 R31599	Chicken nov protel
3	124	15.1	476	1 W53698	Human secreted pro
4	123.5	15.1	425	1 W56778	Homo sapiens D8725
5	123.5	15.1	480	1 W22849	Osteoblast like ce
6	119.5	14.6	258	1 W37466	Inhibitory IGF bin
7	112	13.7	348	1 R25566	Beta-IG-M2. TGF-be
8	112	13.7	348	1 W35731	Rat IGFBP-5. DNA e
9	110.5	13.5	271	1 R26994	Human IGFBP-5. DNA e
10	105	12.8	75	1 R31601	Chicken nov protel
11	109.5	13.4	272	1 R25700	IGFBP6. Insulin-li
12	109.5	13.4	272	1 R26995	Human IGFBP-5. DNA
13	109.5	13.4	272	1 R55084	Human insulin-like
14	109.5	13.4	272	1 R95329	Insulin-like growt
15	109.5	13.4	272	1 W35572	Insulin like growt
16	109	13.3	250	1 W37946	Human connective t
17	110.5	13.5	381	1 W35957	Human monocyte mat
18	103.5	12.6	76	1 R31600	Chicken nov protel
19	109	13.3	347	1 W12694	Connective tissue
20	109	13.3	349	1 R79964	Connective tissue
21	109	13.3	349	1 W11302	Connective tissue
22	109	13.3	349	1 W05089	Human connective t
23	109	13.3	349	1 W62084	Human connective t
24	109	13.3	349	1 W81425	Connective tissue
25	108.5	13.2	381	1 W35730	Human cysteine ric
26	106.5	13.0	258	1 R22253	Sequence of insuli
27	106.5	13.0	258	1 R22253	Sequence of insuli
28	106	12.9	264	1 R89951	Insulin-like growt
29	104	12.7	375	1 R90919	Connective tissue
30	102.5	12.5	291	1 R99952	Insulin-like growt
31	103.5	12.6	379	1 R25565	Beta-IG-M1. TGF-be
32	101.5	12.4	264	1 R13443	FSH inhibiting pro
33	101.5	12.4	264	1 R99950	Recombinant insuli
34	101.5	12.4	264	1 W12343	Human insulin-like
35	101.5	12.4	264	1 W12344	Human insulin-like
36	101.5	12.4	264	1 W12344	Sequence of human
37	101.5	12.4	291	1 R05596	Somatostatin carrie
38	101.5	12.4	291	1 R92300	Insulin like growt
39	101.5	12.4	1122	1 R73954	Human tie tyrosine
40	101.5	12.4	1138	1 R39820	tie receptor kinas
41	103	12.6	1801	1 W50895	Rat laminin B2 cha
42	94.5	11.5	259	1 P91868	Recombinant IGF b1
43	92	11.2	466	1 R07447	Human laminin B1 c

ALIGNMENTS

44	94.5	11.5	1764	1	P91672	Primary amino acid
45	94.5	11.5	1776	1	W50894	Mouse laminin B1 c

ALIGNMENTS

RESULT	1																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																															
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FH Key                                     Location/Qualifiers
FT peptide                               1. 24
FT /label= signal_peptide
FT /note= "only hydrophobic region of protein"
FT binding_site                         56..63
FT /label= IGF-binding_site_motif
FT /note= "corresponds to GCGCCXC consensus"
PN WO9300430-A.
PD 07-JAN-1993.
PE 25-JUN-1992; F00589.
PR 25-JUN-1991; FR-007807.
PA (CNRS ) CENT NAT RECH SCI.
PI Martinerie C, Perbal B;
DR WPI: 93-036377/04.
DR N-PSDB: Q36031.
PT Nucleotide sequences hybridising to regions of chicken nov gene -
PT useful as probes for detecting complementary sequences to
PT evaluate development and/or differentiation of tumours
PS Claim 1; Fig 1; 67pp; French.
CC This amino acid sequence was deduced from the nucleotide sequence
CC of a chicken nov gene clone isolated from a gene bank prepared from
CC chicken embryonic fibroblasts screened with a tumour-derived probe.
CC The only hydrophobic region occurs within the putative signal
CC peptide suggesting that the protein is secreted. The protein also
CC contains the consensus motif of proteins which bind to insulin-like
CC growth factors. It is known that the human IGFII gene is
CC overexpressed in some Wilms' tumours and a similar deregulation of
CC IGFII expression could be involved in nephroblastoma development.
CC The deduced nov protein sequence contains 39 (non-clustered)
CC cysteine residues.
SQ Sequence 351 AA;

Query Match 17.38; Score 141.5; DB 1; Length 351;
Best Local Similarity 31.98; Pred. No. 7.4e-08;
Matches 45; Conservative 17; Mismatches 38; Indels 41; Gaps 7;

QY 3 SVLLTLLVPAHLVAWSNNYAVDCPQHCDSECKSPRC---RTVLDDCGCCRCVCA 59
DB 10 PVLLLLLLLRPCVSGR----EAACPRCGGRCPPAPPCAPGPAVLDDGGCCCLVCAR 65
QY 60 GRGTCYRTVSGMDGKCGPLRCQPSNG-----EDPGEFEFGICK-----DCPYG-- 106
DB 66 QRGES-----CSPLLPCDESGGLYCDRGPD---GGGAGICWLEGDNCVFDGM 111
QY 106 -----TFGMDCRETNCQSG 120
DB 112 IYRNETFPQPSCKYQCTCRDG 132

RESULT 3
W63698
ID W63698 standard; Protein: 476 AA.
AC W63698;
DE Human secreted protein 18.
KW Secreted protein; human; cell proliferation; cytokine activity;
KW tissue growth; cellular differentiation; regeneration; activin;
KW inhibin; chemotactic; haemostatic; thrombolytic; tumour inhibition;
KW anti-inflammatory activity; biomarker.
OS Homo sapiens.
PI Escobedo J, Garcia P, Hu Q, Kothakota S, Williams LT;
PI WPI: 98-348453/30.
DR N-PSDB: V43618.
PT Secreted human polypeptides - having cytokine, cell proliferation or
PT differentiation, activin or inhibin, tumour inhibition or
PT anti-inflammatory activities
PS Claim 1; Pages 70-72; 78pp; English.

This represents a human secreted protein. The specification provides
secreted protein sequences (W63681 to W63699) encoded by the nucleic
acid sequences shown in V43601 to V43619. The invention provides a
method of identifying a secreted polypeptide which is modified by rough
microsomes. The secreted proteins can be used in assays to determine
biological activities, such as cytokine, cell proliferation, or cellular
differentiation activities, tissue growth or regeneration, activin or
inhibin activity, chemotactic or chemokinetic activity, haemostatic or
thrombolytic activity, receptor/ligand activity, tumour inhibition, or
anti-inflammatory activity. The proteins can also be used as biomarkers,
to identify tissues or cell types which express the proteins, or a stage-
or disease-specific alteration in protein expression. They can be used
in protein interaction assays, to identify ligands or binding proteins.
Compounds which affect the biological activities of the secreted proteins
or their ability to interact with specific ligands can be identified
using the proteins in screening assays. The proteins and antibodies that
bind specifically to the protein can also be used to design diagnostic
tests and therapeutic compositions for diseases which may be associated
with altered expression of these proteins. Fusion proteins comprising,
e.g. signal sequences or transmembrane domains of the proteins can be
used to target other protein domains to cellular membrane or they can
be secreted extracellularly.
SQ Sequence 476 AA;

Query Match 15.18; Score 124; DB 1; Length 476;
Best Local Similarity 40.08; Pred. No. 7e-06;
Matches 34; Conservative 13; Mismatches 28; Indels 10; Gaps 4;

QY 5 LLTLLTLLVPAHLVAWSNNYAVDCPQHCDSECKSPRC---KRTVLDDCGCCRCVCAAGR 61
DB 19 LLL--LLVPVWAGAEKLHTQPCPVCQPTRCPLTTCALGTTPVFDLCRCRCVCPRAE 76
QY 62 GETCYRTVSGMDGKCGPLRC-OP 85
DB 77 REVC---GGAQGPCAPGLQLQP 97

RESULT 4
W56778
ID W56778 standard; Protein: 425 AA.
AC W56778;
DE Homo sapiens D87258 sequence.
KW PS-1; Presenilin; presenilin-1; PSP-1; Alzheimer's disease;
KW serine protease; neurodegeneration; predisposition; diagnosis;
KW D87258.
OS Homo sapiens.
PI Key Location/Qualifiers
FT Misc_difference 213 /label= Gly, Val
FT EP-828003-A2.
PN 11-MAR-1998.
PD 26-AUG-1997; 306501.
PR 13-DEC-1996; US-032875.
PR 06-SEP-1996; US-025436.
PR 25-OCT-1996; US-027873.
PA (SMIK ) SMITHKLINE BEECHAM CORP.
PA (SMIK ) SMITHKLINE BEECHAM PLC.
PI Browne MJ, Clankinbeard HE, Creasy CL, Karran EH,
PI WPI: 98-161101/15.
DR N-PSDB: V29540.
PT Nucleic acids encoding human serum protease protein(s) - used for
PT diagnosing pre-disposition to Alzheimer's disease, etc.
PS Claim 22; Page 30-31; 65pp; English.
CC The sequence is that encoded by cDNA clone D87528 which can be used
CC to identify modulators of serine protease activity and also to diagnose
CC a condition associated with lack of one of the serine proteases
CC or a genetic predisposition to neurodegeneration in a patient,
CC preferably predisposition to Alzheimer's disease.
SQ Sequence 425 AA;

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RESULT 12
26995

RESULT 13

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R55084      R55084 standard; Protein; 272 AA.
ID          ID
AC          AC
R55084;    R55084;
28-NOV-1994 (first entry)
DE          DE
Human insulin-like growth factor binding protein-5.
DD         DD
Insulin-like growth factor binding protein-5; IGFBP-5; hormone.
OS         OS
Homo sapiens.
Key        Key
EFH        EFH
peptide    peptide
Location/Qualifiers
24..163
/note= "preferred truncated IGFBP-5, Claim 12"
EF         EF
peptide    peptide
21..189
/note= "preferred truncated IGFBP-5, Claim 13"
EF         EF
WO9410207-A.
MN         MN
11-MAY-1994.
PD         PD
29-OCT-1993; U10462.
FF         FF
04-NOV-1992; US-972142.
RR         RR
(CHIR ) CHIRON CORP.
CA         CA
Address DL, Kiefer MC;
IL         IL
WPI; 94-167395/20.
RR         RR
N-PSDB; Q65519.
RR         RR
Truncated insulin-like growth factor binding protein - has
reduced affinity for insulin-like growth factor, useful for
stimulating bone cell growth and mitogenic activity
Disclosure; Fig. 1; 56pp; English.
This protein sequence are contained 2 preferred truncated

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CC IGFBP-5 proteins derived from human U-2 osteosarcoma cells.
CC The truncated IGFBPs may be used for stimulating mitogenic
CC activity, particularly for stimulating bone cell growth. The
CC IGFBP is preferably produced recombinantly by expression in a
CC yeast or CHO host.
SQ Sequence 272 AA;
SQ

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Query Match      13.4%; Score 109.5; DB 1; Length 272;
Best Local Similarity 27.7%; Pred. No. 0.00014;
Matches         26; Conservative 20; Mismatches 27; Indels 21; Gaps 3;
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QY 4 VLLTTLVPAHLVAANSNNVADCPQHCDSECKSPRCRKTWLD-----DCGCCRV 56
:| |: | |: | |: | |: | |: | |: |
Db 8 LLLLAAYAGPAAQSLSGFV-----HCEPCDEKALSMCPPSLGCELNVKEPGGCMT 58

QY 57 CAGRGTCCTRTVSGMDGMKGCGPELRCQPNSNGED 90
|| :| |: | |: | |: | |: | |: |
Db 59 CALAEGOSC-----GVYTERCAOGRCLPRODEF 87

RESULT	14	
R95329		
ID	R95329 standard; Protein; 272 AA.	
AC	R95329;	
DE	09-DEC-1996 (first entry)	
DT	Insulin-like growth factor binding protein-5 (IGFBP-5).	
DE	Insulin-like growth factor binding protein-5 (IGFBP-5).	
KW	TNF-R1-PD; tumour necrosis factor receptor 1 death domain; inhibitor;	
KW	P55; anti-inflammatory; autoimmune disease; graft versus host reaction;	
KW	osteoporosis; cachexia; diabetes; sequence identity; IGFBP-5;	
KW	insulin-like growth factor binding protein-5.	
OS	Homo sapiens.	
FT	Key	Location/Qualifiers
FT	protein	87..272
FT		/label= clone 20DD

PN	W09612735-A1.
PD	12-MAY-1996.
PD	12-OCT-1995: U12724.
PF	19-OCT-1994; US-327514.
PR	19-JUN-1995; US-494440.
PR	26-SEP-1995; US-533901.
PR	(GENY) GENETICS INST INC.
PA	Chen J, Graham J, Lin L, Schievella AR;
PI	WPI; 96-230551/23.
DR	N-PSDB; T15231.
DR	TNF receptor death domain ligand proteins and inhibitors of ligand
PT	binding - for prevention and treatment of pref. anti-inflammatory
PT	conditions, e.g. auto-immune disease, graft versus host reaction
PT	osteoporosis, etc.
PT	Claim 15; Page 42-43; 83pp; English.
PS	The present sequence is that of insulin-like growth factor binding
CC	protein-5 (IGFBP-5). Based upon the amino acid sequence identity between
CC	IGFBP-5 and a tumour necrosis factor (TNF) receptor 1 (R1) death domain
CC	(DD) ligand (clone 20DD; R95328) it has been determined that IGFBP-5 and
CC	certain fragments of it, will exhibit TNF-R1-DD ligand binding activity.
CC	A yeast genetic selection method, the "interaction trap", was used to
CC	screen W138 cell cDNA libraries for proteins that interact with the DD
CC	of the p55 type 1 TNF-R. TNF-R1-DD ligands and their inhibitors, e.g.
CC	IGFBP-5, are useful in the prevention and treatment of anti-inflammatory
CC	conditions and other conditions such as cachexia, autoimmune disease,
CC	graft versus host reaction, osteoporosis, diabetes, etc.
CC	Sequence 272 AA;
SQ	

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Query Match      13.48; Score 109.5; DB 1; Length 272;
Best Local Similarity 27.7%; Pred. No. 0.00014;
Matches 26; Conservative 27; Mismatches 21; Gaps 3;

QY   4 VLLLTTLVPAHLVAASNNYAVDCPQHDSSECKSPRCKRTVLDP-----DCGCCRV 56
      |||||  A|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|::|
Db    8 LLLLLAAAYGAQSLGSGFY-----HCFPCDEKALSMCPPPLGCGLVKPEPGCCMT 58
      e

QY   57 CAAGRGETCYRTVSGMDGMKGCPGLRCOPNGED 90

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Db      || : |:| | : |: |:| | | | :
59 CALAEQSC-----GVYTERCAQGLRCLPRQDEE 87
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RESULT 15	
W35572	ID W35572 standard; Protein; 272 AA.
AC	W35572;
DT	19-MAR-1998 (first entry)
DE	Insulin like growth factor 5 binding protein.
DE	Insulin like growth factor receptor P55 type; TNF-R1-DD ligand protein;
KW	Tumour necrosis factor
KW	death domain; TNF-R1; inhibitor identification; TNF induced condition;
KW	insulin-like growth factor binding protein-5; inflammatory condition;
KW	IGFBP-5; therapy.
OS	Homo sapiens.

PN	WO9730084-A1.
PD	21-AUG-1997.
PF	11-FEB-1997; U02146.
PR	15-AUG-1996; US-698551.
PR	15-FEB-1996; US-602228.
PA	(GEMY) GENETICS INST INC.
PI	Chen J, Graham J, Lin L, Schievella AR;
DR	WPI; 97-424976/39.
DR	N-PSDB; T94634.
PT	Tumour necrosis factor receptor p55 type death domain ligand
PT	proteins - useful for preventing or ameliorating inflammatory
PT	conditions
PS	Claim 15; Page 45-46; 103pp; English.
CC	This sequence represents a protein sequence of the invention. This
CC	sequence is the insulin-like growth factor binding protein-5 (IGFBP-5)
CC	and is a tumour necrosis factor receptor p55 type (TNF-R1) death domain
CC	(DD) ligand protein. A host cell containing DNA encoding this sequence is
CC	used for the recombinant production of TNF-R1-DD. The TNF-R1-DD ligand
CC	protein can be used in a method to identify inhibitors of TNF-R DD
CC	binding. The TNF-R1-DD ligand protein, IGFBP-5 (has TNF-R1-DD ligand
CC	activity), or inhibitors of TNF-R1-DD ligand protein are capable of
CC	preventing or ameliorating an inflammatory condition, preferably by
CC	inhibiting TNF-R DD binding. Identification and isolation of ligands
CC	allows their effects upon TNF-R signal transduction and use as
CC	therapeutic agents for treatment of TNF-induced conditions to be
CC	examined.
CC	Sequence 272 AA:
SQ	Sequence

Query Match	13.4%	Score 109.5;	DB 1;	Length 272;
Best Local Similarity	27.7%;	Pred. No. 0.00014;		
Matches 26; Conservative	20;	Mismatches 27;	Indels 21;	Gaps 3

	Qy	4	VLLTTLVPAILVAHNSNNVAVDCPOHCSSSCKSPKCRKTVD-----DGCGRV	56
			: : : : : :	
	Db	8	LIIILAAAGPAQSGLSFV-----HCPEDEKALSMCPPSLGCELVKEPGGCMT	58
			: : : : : :	
	Qy	57	CAARGETCYRTVSMDGMKGPGRLRQPNGED	90
			: : : : : :	
	Db	59	CALAIGQSC---GVYTERCAOGLCLPRODEE	87
			: : : : : :	

Search completed: May 4, 1999, 12:32:37
Job time: 9784 sec

Primary amino acid
Mouse laminin B1 c

ALIGNMENTS

44	94.5	10.3	1764	1	P91672
45	94.5	10.3	1776	1	W50894

AC	R39394;
DT	06-NOV-1996 (first entry)
DE	Vascular IBP-like growth factor.
DE	Vascular IBP-like growth factor.
KW	Insulin-like growth factor binding protein; VIGF;
KW	muscle wastage; osteoporosis; implant fixation; wound healing;
KW	therapv; diagnosis.

growth factor binding

therapy; diagnosis
Homo sapiens

Location/Qualifiers
1. 21

PT	WO9617931-A1.
PN	13-JUN-1996.
PD	09-DEC-1994; U14388.
PF	09-DEC-1994; WO-U14388.
PA	(HUMA-) HUMAN GENOME SCI INC.
PR	Hastings GA, Rosen CA;
PI	WPI; 96-287176/29.
DR	N-PSDB; T34991.
PT	Human vascular insulin-like growth factor binding protein-like
PT	growth factor, and its nucleic acid sequence and (antagonists -
PT	used, e.g. to treat muscle wasting diseases or aid implant fixation,
PT	or limit excess connective tissue prodn. during wound healing.
PS	Claim 14; Page 43-44; 61pp; English.
CC	Human vascular insulin-like growth factor binding protein-like
CC	growth factor (R98994), or VIGF, is a protein of primarily
CC	vascular origin that is structurally related to the IBP and CCN
CC	protein families. It can be expressed in e.g. E. coli, CHO or
CC	insect host cells using a vector incorporating a cDNA clone
CC	(T34991), or its derivative, obtd. from human umbilical
CC	endothelial cells. It is useful therapeutically e.g. for
CC	treating muscle wasting diseases or osteoporosis, or can be used
CC	to detect diseases associated with under- or over-expression of VIGF,
CC	or to screen for antagonists useful during wound healing.
SO	Sequence 184 AA:

WO-014388. T GENOME SCT INC

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Query Match          100.0%; Score 918; DB 1; Length 184;
Best Local Similarity 100.0%; Pred. NO. 2.7e-94;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  MKSVLLTLLVPAHLVAWSNNYAVDCPOHCDSDSECKSSPRCKRTVLDDCGCCRVCAAG  60
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db      1  MKSVLLTLLVPAHLVAWSNNYAVDCPOHCDSDSECKSSPRCKRTVLDDCGCCRVCAAG  60

QY      61  RGEFCYRTVSGMDGKCGPLRCOPSGNEDPFGEEFGICKDCPYGTGMDCRETNCQSG  120
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db      61  RGEFCYRTVSGMDGKCGPLRCOPSGNEDPFGEEFGICKDCPYGTGMDCRETNCQSG  120

QY      121  ICDRGTKCLKFFPFQYSVTKSSNRFVSLTEHDMASGDGNIVREEVVKENAAAGSPVMRKW  180
      | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db      121  ICDRGTKCLKFFPFQYSVTKSSNRFVSLTEHDMASGDGNIVREEVVKENAAAGSPVMRKW  180

QY      181  LNPR 184
      | | | |
Db      181  LNPR 184

RESULT      2
R31599
ID      R31599 standard; Protein; 351 AA.
AC      R31599;
DT      24-MAY-1993 (first entry)

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DE Chicken nov protein.
 KW avian nephroblastoma; avian myeloblastoma virus; IGF binding site;
 OS insulin-like growth factor; Wilm's tumour.
 FH Gallus domesticus.
 Key Location/Qualifiers
 FT peptide 1..24
 FT /label= signal_peptide
 FT /note= "only hydrophobic region of protein"
 FT binding_site 56..63
 FT /label= IGF-binding_site_motif
 FT /note= "corresponds to GCGCCXC consensus"
 PN W03900430-A.
 FN 07-JAN-1993.
 PD 25-JUN-1992; F00589.
 PE 25-JUN-1991; FR-007807.
 PR (CNRS) CENT NAT RECH SCI.
 PI Martinerie C, Perbal B;
 DR WPI; 93-036377/04.
 DR N-PSDB; Q36031.
 PT Nucleotide sequences hybridising to regions of chicken nov gene -
 useful as probes for detecting complementary sequences to
 evaluate development and/or differentiation of tumours
 PS Claim 1; Fig 1; 67pp; French.
 CC This amino acid sequence was deduced from the nucleotide sequence
 of a chicken nov gene clone isolated from a gene bank prepared from
 chicken embryonic fibroblasts screened with a tumour-derived probe.
 CC The only hydrophobic region occurs within the putative signal
 peptide suggesting that the protein is secreted. The protein also
 contains the consensus motif of proteins which bind to insulin-like
 growth factors. It is known that the human IGFII gene is
 overexpressed in some Wilm's tumours and a similar deregulation of
 IGFII expression could be involved in nephroblastoma development.
 CC The deduced nov protein sequence contains 39 (non-clustered)
 cysteine residues.
 SQ Sequence 351 AA;

Query Match 15.4%; Score 141.5; DB 1; Length 351;
 Best Local Similarity 31.9%; Pred. No. 3e-08;
 Matches 45; Conservative 17; Mismatches 38; Indels 41; Gaps 7;

QY 3 SVLLLTLLVPAHLVAWSNNYAVDCPQHCDSECKSSPRCK---RTVLDDCGCCRVCA 59
 DB 10 PVLLLLLLLLRCVSGR---EACPRCGGRCAPGVPVAVLDGCGCLVCAR 65
 QY 60 GRGETCYRTVSGMDGKMGKGLRCQPSNG-----EDPFGEEFGICK-----DCPYG-- 106
 DB 66 QRGES-----CSPLLPCDESGLYCDRGPED--GGGAGICWLEGDNCVFDGM 111
 QY 106 -----TFGMDCRETCNCQSG 120
 DB 112 IYRNETFPQSKYQCTCRDG 132

RESULT 3
 ID W63698 standard; Protein: 476 AA.
 AC W63698.
 DT 24-SEP-1998 (first entry)
 DE Human secreted protein 18.
 KW Secreted protein; human; cell proliferation; cytokine activity;
 KW tissue growth; cellular differentiation; regeneration; activin;
 KW inhibin; chemotactic; haemostatic; thrombolytic; tumour inhibition;
 KW anti-inflammatory activity; biomarker.
 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT Misc_difference 213
 FT /label= Gly, Val
 PN EP-828003-A2.
 PD 11-MAR-1998.
 PE 26-AUG-1997; 306501.
 PR 13-DEC-1996; US-032875.
 PR 06-SEP-1996; US-025436.
 PR 25-OCT-1996; US-027873.
 PA (SMIK) SMITHKLINE BEECHAM CORP.
 PA (SMIK) SMITHKLINE BEECHAM PLC.
 PI Browne MJ, Clinkenbeard HE, Creasy CL, Karran EH,
 PI Livi GP, Southan CD;
 DR WPI; 98-161101/15.
 DR N-PSDB; V29540.
 PT Nucleic acids encoding human serum protease protein(s) - used for
 diagnosing pre-disposition to Alzheimer's disease, etc.
 PS Claim 22; Page 30-31; 65pp; English.
 CC The sequence is that encoded by cDNA clone D87528 which can be used
 to identify modulators of serine protease activity and also to diagnose
 a condition associated with lack of one of the serine proteases

PT Secreted human polypeptides - having cytokine, cell proliferation or
 differentiation, activin or inhibin, tumour inhibition or
 anti-inflammatory activities
 PS Claim 1; Pages 70-72; 78pp; English.
 CC This represents a human secreted protein. The specification provides
 secreted protein sequences (W63681 to W63699) encoded by the nucleic
 acid sequences shown in V43601 to V43619. The invention provides a
 method of identifying a secreted polypeptide which is modified by rough
 microsome. The secreted proteins can be used in assays to determine
 biological activities, such as cytokine, cell proliferation, or cellular
 differentiation activities, tissue growth or regeneration, activin or
 inhibin activity, chemotactic or chemokinetic activity, haemostatic or
 thrombolytic activity, receptor/ligand activity, tumour inhibition, or
 anti-inflammatory activity. The proteins can also be used as biomarkers,
 or to identify tissues or cell types which express the proteins, or a stage-
 or disease-specific alteration in protein expression. They can be used
 in protein interaction assays, to identify ligands or binding proteins.
 CC Compounds which affect the biological activities of the secreted proteins
 or their ability to interact with specific ligands can be identified
 using the proteins in screening assays. The proteins and antibodies that
 bind specifically to the protein can also be used to design diagnostic
 tests and therapeutic compositions for diseases which may be associated
 with altered expression of these proteins. Fusion proteins comprising,
 e.g. signal sequences or transmembrane domains of the proteins can be
 used to target other protein domains to cellular membrane or they can
 be secreted extracellularly.
 SQ Sequence 476 AA;

Query Match 13.5%; Score 124; DB 1; Length 476;
 Best Local Similarity 40.0%; Pred. No. 3.5e-06;
 Matches 34; Conservative 13; Mismatches 28; Indels 10; Gaps 4;

QY 5 LLLTLLVPAHLVAWSNNYAVDCPQHCDSECKSSPRCK---KRTVLDDCGCCRVCA 61
 DB 19 LLL--LLVPVLWAGAEKLTQPCPAVCQPTRCPALPTCAIGTTPVFDLCRCRCVCPAAE 76
 QY 62 GETCYRTVSGMDGKMGKGLRC-QP 85
 DB 77 REVG-----GGAQGQPCAPGLQCLQP 97

RESULT 4
 W56778
 ID W56778 standard; Protein: 425 AA.
 AC W56778.
 DT 13-OCT-1998 (first entry)
 DE Homo sapiens D87258 sequence.
 KW PS-1; presenilin; presenilin-1; PSP-1; Alzheimer's disease;
 KW serine protease; neurodegeneration; predisposition; diagnosis;
 KW D87258.
 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT Misc_difference 213
 FT /label= Gly, Val
 PN EP-828003-A2.
 PD 11-MAR-1998.
 PE 26-AUG-1997; 306501.
 PR 13-DEC-1996; US-032875.
 PR 06-SEP-1996; US-025436.
 PR 25-OCT-1996; US-027873.
 PA (SMIK) SMITHKLINE BEECHAM CORP.
 PA (SMIK) SMITHKLINE BEECHAM PLC.
 PI Browne MJ, Clinkenbeard HE, Creasy CL, Karran EH,
 PI Livi GP, Southan CD;
 DR WPI; 98-161101/15.
 DR N-PSDB; V29540.
 PT Nucleic acids encoding human serum protease protein(s) - used for
 diagnosing pre-disposition to Alzheimer's disease, etc.
 PS Claim 22; Page 30-31; 65pp; English.
 CC The sequence is that encoded by cDNA clone D87528 which can be used
 to identify modulators of serine protease activity and also to diagnose
 a condition associated with lack of one of the serine proteases

CC or a genetic predisposition to neurodegeneration in a patient,
CC preferably predisposition to Alzheimer's disease.
SQ Sequence 425 AA;

Query Match 13.5%; Score 123.5; DB 1; Length 425;
Best Local Similarity 34.0%; Pred. No. 3.6e-06;
Matches 33; Conservative 18; Mismatches 25; Indels 21; Gaps 5;
QY 4 VLLTLLVPAHLV-RAWSNNYAVDCPOHDSSECKSSP-----RCKRTVLDDCGCCRV 56
DB 12 LLLLLAAPASQUSRAPLAAGCPDRCEPARCPQPEHCEGGRAR----DAGCCCEV 67
QY 57 CAAGGTCYRTVSGMDGMKCGPLRCQPSNGEDPFG 93
DB 68 CGAPEGAAAC-----GLQEGPCGEGLCQV-----PFG 94

RESULT 5
W22849 ID W22849 standard; Protein; 480 AA.
AC W22849;
DT 16-SEP-1997 (first entry)
DE Osteoblast like cell derived protein.
KW Osteoblast like cell; prevention; treatment; disease;
OS Homo sapiens.
FH Key Location/Qualifiers
FT peptide 1..27
FT peptide /label= sig_peptide
FT peptide 28..480
FT peptide /label= mat_peptide
PN J09107980-A.
PD 28-APR-1997.
PF 12-AUG-1996; 231415.
PR 17-AUG-1995; JP-233537.
PA (NIBS) JAPAN TOBACCO INC.
DR WPI; 97-292469/27.
DR N-PSDB; T75444.
PT DNA encoding osteoblast like cell derived protein - useful for treatment and prevention of various diseases and as contraceptive
PS Claim 2; Pages 26-27; 42pp; Japanese.
CC The present sequence is an osteoblast like cell derived protein, which may be used to prevent or treat various diseases, or as a contraceptive.
CC mRNA was collected from an osteoblast like cell, and double stranded DNA synthesised. A 3'-directed cDNA library was prepared, and the base sequence of the DNA of each clone in the cDNA library determined. An osteoblast complete chain cDNA library was prepared, and the clone GS2422, which has a complete chain, isolated. GS2422 protein was expressed and purified, and an antibody against GS2422 prepared. The amino acid sequence of a purified specimen of GS2422 protein was analysed. The GS2422 gene is expressed in various human organs.
SQ Sequence 480 AA;

Query Match 13.5%; Score 123.5; DB 1; Length 480;
Best Local Similarity 34.0%; Pred. No. 4.1e-06;
Matches 33; Conservative 18; Mismatches 25; Indels 21; Gaps 5;
QY 4 VLLTLLVPAHLV-RAWSNNYAVDCPOHDSSECKSSP-----RCKRTVLDDCGCCRV 56
DB 12 LLLLLAAPASQUSRAPLAAGCPDRCEPARCPQPEHCEGGRAR----DAGCCCEV 67
QY 57 CAAGGTCYRTVSGMDGMKCGPLRCQPSNGEDPFG 93
DB 68 CGAPEGAAAC-----GLQEGPCGEGLCQV-----PFG 94
RESULT 6
W37466 ID W37466 standard; Protein; 258 AA.

AC W37466;
DT 28-MAY-1998 (first entry)
DE Inhibitory IGF binding protein.
KW Inhibitory insulin like growth factor binding protein; In-IGF-BP;
OS Insulin like growth factor; IGF; Bone cell; bone tumour; inhibition.
PN Homo sapiens.
PD US5693754-A.
PD 02-DEC-1997.
PF 22-OCT-1992; 966121.
PR 03-JUL-1990; US-548388.
PR 22-OCT-1992; US-966121.
PA (BOEF) BOEHRINGER MANNHEIM GMBH.
PI Baylink DJ, Mohan S;
DR WPI; 98-031813/03.
DR N-PSDB; V00257.
PT Inhibitory insulin like growth factor binding protein - useful to inhibit effects of insulin like growth factor on bone cells and in diagnostic assays
PS Claim 1; Fig 1; lpp; English.
CC This amino acid sequence is the novel inhibitory insulin like growth factor binding protein (In-IGF-BP), which is 258 amino acid residues in length. The In-IGF-BP can be used to modulate the effects of insulin like growth factor I or II (IGF) on bone cells. It is used to treat IGF dependent bone tumours, and in a competitive binding assay to determine IGF in clinical samples.
SQ Sequence 258 AA;

Query Match 13.0%; Score 119.5; DB 1; Length 258;
Best Local Similarity 32.4%; Pred. No. 6e-06;
Matches 34; Conservative 24; Mismatches 30; Indels 17; Gaps 6;
QY 6 LLTTLVPAHLVAAWSNNYAVDCPOHCD---SSECKSSPCKRTVLDDCGCCRVCAARG 62
DB 6 LVAALLAAGPGLSLGDE-AIHCPGCEKLAGRCPPVGCCELVRACGCCATCALGLG 64
QY 63 ETCYRTVSGMDGMKCGPLRCQPSNG-EDPF-----GEEFGICKD 101
DB 65 MPC-----GVYTPRCGSLRCYPPRGVEXPLHTLMHGE--GVCME 102

RESULT 7
R25566 ID R25566 standard; Protein; 348 AA.
AC R25566;
DT 18-JAN-1993 (first entry)
DE Beta-IG-M2.
KW Transforming growth factor beta; induced; CEF-10; v-src; chicken;
KW embryo; fibroblasts; TGF-beta.
OS Mus musculus.
PN EP-495674-A.
PD 22-JUL-1992.
PF 17-JAN-1992; 300429.
PR 18-JAN-1991; US-642991.
PR 10-JAN-1992; US-816270.
PA (BRIM) BRISTOL-MYERS SQUIBB CO.
PI Brunner AM, Chinn J, Neubauer MG, Purchio AF;
DR WPI; 92-243508/30.
DR N-PSDB; Q26422.
PT TGF-beta induced gene family - encodes proteins involved in growth and differentiation effects of TGF-beta-1
PS Claim 3; Fig 2; 35pp; English.
CC The protein sequence was deduced from the DNA sequence obtd. by screening a cDNA library made from AKR-2B mouse cells induced with TGF-beta1 and cyclohexamide with two probes from untreated AKR-2B mRNA and AKR-2B mRNA from cells treated with cyclohexamide and TGF-beta1. The proteins encoded by hybridising colonies (beta-IG-M1 and beta-IG-M2) contain 38 Cys residues and are induced by TGF-beta1.
CC Beta-IG-M2 displays 50 percent homology to the CEF-10 protein induced by v-src in chicken embryo fibroblasts. Residues 52-59 of beta-IG-M2 conform to the GCGCCXCC motif reported in the amino half of insulin-like growth factor (IGF) binding proteins.
CC The C-terminal Cys rich region of beta-IG-M1, -M2 and CEF-10 contain

an amino acid sequence with strong homology to a motif found near the C-terminal of the malarial circumsporozoite (CS) protein, which is highly conserved among all species of malarial parasites sequenced to date (designated region II). This motif is also found in other proteins which have cell adhesive properties that mediate cell-cell and cell-extracellular matrix interactions, such as properdin, thrombospondin, and TRAP. The proteins encoded by TGF-beta induced genes are likely to be involved in mediation of the biological effects of TGF-beta relating to cell growth and differentiation. See also R25565.

Sequence 348 AA;
Query Match 12.2%; Score 112; DB 1; Length 348;
Best Local Similarity 30.4%; Pred. No. 5.4e-05;
Matches 34; Conservative 15; Mismatches 35; Indels 28; Gaps 7;

QY 27 DCPQHDSSECKSSPRCK---RTVLDDCGCCRCVCAAGRGCTCYRTVSGMDGMKCGP--GL 81
DB 27 DCSAQCCCAA-EAAPHCPAGVSLVLDGCGCCRCVCAKQGLGELC-----TERDPCDPHKL 79
QY 82 RCQPSNGEDPGEFFGICKD-----CPVG-----TFGMDCRETNCQSG 120
DB 80 FCDFGS---PANRIGVCTAKDGAPCVFGSVYRSGESFQSSCKYQCTCLDG 128

RESULT 8
W35731 ID W35731 standard; Protein; 348 AA.
AC W35731; 27-MAR-1998 (first entry)
DE Murine Fisp12
KW Fisp12; cysteine rich protein; mouse; Cyr61;
KW extracellular matrix signalling molecule; cell adhesion;
KW cell migration; cell proliferation; angiogenesis; chondrogenesis;
KW oncogenesis.
OS Mus musculus.
PN W09733995-A2.
PD 18-SEP-1997; U04193.
PF 14-MAR-1997; U04193.
PR 15-MAR-1996; US-013958.
PA (MUN1-) MUNIN CORP.
PI Lau LF;
DR WPI: 97-470875/43.
DR N-PSDB; T94700.
PT Isolated and purified cysteine rich protein 61, Cyr61 - useful to modulate e.g. haematostasis, induce wound healing, promote organ regeneration etc
PT Example 2; Page 115-116; 133pp; English.
PS This protein sequence comprises murine Fisp12, an extracellular matrix signalling molecule (ECM) that exhibits structural similarity to Cyr61 (see W35730) and which, like Cyr61, influences cell adhesion, proliferation and migration. The human orthologue of Fisp12 is connective tissue growth factor. Fisp12 polynucleotides (see T94700) can be used for the production of Fisp12 polypeptides by recombinant methods. Polypeptide compositions are provided that comprise mammalian ECM signalling molecules, peptide fragments, inhibitory peptides capable of interacting with receptors for ECM signalling molecules, and antibody products. Further provided are methods for using mammalian ECM signalling molecules to screen for, and/or modulate disorders associated with angiogenesis, chondrogenesis and oncogenesis; ex vivo methods for using ECM signalling molecules to prepare blood products are also provided.

Query Match 12.2%; Score 112; DB 1; Length 348;
Best Local Similarity 30.4%; Pred. No. 5.4e-05;
Matches 34; Conservative 15; Mismatches 35; Indels 28; Gaps 7;
QY 27 DCPQHDSSECKSSPRCK---RTVLDDCGCCRCVCAAGRGCTCYRTVSGMDGMKCGP--GL 81
DB 27 DCSAQCCCAA-EAAPHCPAGVSLVLDGCGCCRCVCAKQGLGELC-----TERDPCDPHKL 79

Db 27 DCSAQCCCAA-EAAPHCPAGVSLVLDGCGCCRCVCAKQGLGELC-----TERDPCDPHKL 79
QY 82 RCQPSNGEDPGEFFGICKD-----CPVG-----TFGMDCRETNCQSG 120
DB 80 FCDFGS---PANRIGVCTAKDGAPCVFGSVYRSGESFQSSCKYQCTCLDG 128
RESULT 9
R26994 ID R26994 standard; Protein; 271 AA.
AC R26994;
DT 16-FEB-1993 (first entry)
DE Rat IGFBP-5.
KW rat insulin-like growth factor binding protein-5; IGF-I; IGF-II;
KW breast cancer; bone cancer; modulating bone growth; purification;
KW affinity columns; antibodies; diagnosis; testing; ss.
OS Synthetic.
FH Key Location/Qualifiers
FT peptide i.19 /note= "signal peptide"
PN W09214834-A.
PD 03-SEP-1992.
PF 13-FEB-1992; U01196.
PR 14-FEB-1991; US-658410.
PA (WHIT-) WHITTIER INST DIABETES & ENDOCRINOLOGY.
PI Ling NC, Shimaski S;
DR WPI: 92-316186/38.
DR N-PSDB; Q28270.
PT DNA encoding insulin-like growth factor binding protein - useful for treating breast and bone cancer and modulating bone growth
PS Disclosure; Page 11; 42pp; English.
CC This sequence represents insulin-like growth factor binding protein. (IGBP-5) It was deduced from the appropriate nucleotide sequence obtained as in Q28270. IGBP-5 can bind to both IGFs -I and -II. It is useful for treating conditions caused by an overabundance of IGFs eg. certain breast cancers and bone cancers. It is useful for modulation of bone growth and in affinity chromatography columns for purification of IGF -I and -II. Antibodies to it can be used in assays to detect levels of the protein in mammals esp. humans and to neutralise the effects of IGFBP-5, and are useful in diagnostic test kits.
SQ Sequence 271 AA;

Query Match 12.0%; Score 110.5; DB 1; Length 271;
Best Local Similarity 28.7%; Pred. No. 6.2e-05;
Matches 27; Conservative 20; Mismatches 26; Indels 21; Gaps 3;
QY 4 VLLTTLLVPAHLVAWSNNYAVDCPHQDSECKSSPRCKRTVLD-----DCGCCRV 56
DB 7 LLLLAACAVPAQGLGSFV-----HCEPCDEKALSMCPPSPGLGCELVKPGCGCMT 57
QY 57 CAAGRGCTCYRTVSGMDGMKCGPGLRCQPSNGED 90
DB 58 CALAESQSC-----GVYTERCAQGLRCLPRQDEE 86

RESULT 10
R31601 ID R31601 standard; Protein; 75 AA.
AC R31601;
DT 24-MAY-1993 (first entry)
DE Chicken nov protein fragment V.
KW avian nephroblastoma; avian myeloblastoma virus;
KW stringent hybridisation.
OS Gallus domesticus.
PN W09300430-A.
PD 07-JAN-1993.
PF 25-JUN-1992; F00589.
PR 25-JUN-1991; FR-007807.
PA (CNRS) CENT NAT RECH SCI.
PI Martinerie C, Perbal B.
DR WPI: 93-036377/04.
PT Nucleotide sequences hybridising to regions of chicken nov gene -

RESULT	13		
R55084			
ID	R55084	standard; Protein; 272 AA.	
AC	R55084;		
DT	28-NOV-1994	(first entry)	
DE	Human insulin-like growth factor binding protein-5.		
KW	Insulin-like growth factor binding protein-5; IGFBP-5; hormone.		
OS	Homo sapiens.		
EH	Key	Location/Qualifiers	
FT	peptide	24..163	
FT	peptide	/note="preferred truncated IGFBP-5, Claim 12"	
FT	peptide	21..189	
FT	peptide	/note="preferred truncated IGFBP-5, Claim 13"	
PN	WO9410207-A.		
PD	11-MAY-1994.		
PF	29-OCT-1993;	U10462.	
PR	04-NOV-1992;	US-972142.	
PA	(CHIR) CHIRON CORP.		
PI	Address DL, Kiefer MC;		
DR	WPI; 94-167395/20.		
DR	N-PGDB; Q65519.		
PT	Truncated insulin-like growth factor binding protein - has		


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DR EMBL; U02025; G437125; ..
DR EMBL; U02023; G437125; JOINED.
DR EMBL; U02027; G437125; JOINED.
DR EMBL; U02024; G437125; JOINED.
DR EMBL; X81583; G550385; -.
DR MGD; MGI:96440; IGFBP5.
DR PROSITE; PS00222; IGF-BINDING; 1.
DR PROSITE; PS00484; THYROGLOBULIN_1; 1.
KW GROWTH FACTOR BINDING; SIGNAL.
FT SIGNAL 1 19 POTENTIAL.
FT CHAIN 20 271 INSULIN-LIKE GROWTH FACTOR BINDING
FT DOMAIN 214 262 THYROGLOBULIN TYPE 1.
FT CONFLICT 112 112 MISSING (IN REF. 2).
SQ SEQUENCE 271 AA; 30372 MW; 12DC64CA CRC32;

Query Match 12.5%; Score 114.5; DB 1; Length 271;
Best Local Similarity 28.7%; Pred. No. 2.1e-05;
Matches 27; Conservative 20; Mismatches 26; Indels 21; Gaps 3;

QY 4 VLLITLLVPAHLVAANSNYAVDCPHQDSSECKSSPRCKRTVLD-----DCGCCR 56
DB 7 LLLLAAYAVPAQGLGSEV-----HCEPCDEKALSMCPPSPGLGELVKPEPGCCMT 57
QY 57 CAAGRGTCYRTVSGDMGKPGCLRCOPSGED 90
DB 58 CALAEGGSC-----GVYTERCAQGLRCLPRODEE 86

RESULT 7
CTGF_MOUSE
ID CTGF_MOUSE STANDARD; PRT; 348 AA.
AC P29268;
DT 01-DEC-1992 (REL. 24, CREATED)
DT 01-DEC-1992 (REL. 24, LAST SEQUENCE UPDATE)
DE 15-JUL-1998 (REL. 36, LAST ANNOTATION UPDATE)
DE CONNECTIVE TISSUE GROWTH FACTOR PRECURSOR (CTGF) (FISP-12 PROTEIN).
GN CTGF OR FISP12 OR FISP-12.
OS MUS MUSCULUS (MOUSE).
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;
OC EUTHERIA; RODENTIA.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 91363290.
RA RYSECK R.-P., MACDONALD-BRAVO H., MATTEI M.-G., BRAVO R.;
RL CELL GROWTH DIFFER. 2:225-233(1991).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE; 91229699.
RA BRUNNER A., CHINN J., NEUBAUER M.G., PURCHIO A.F.;
RL DNA CELL BIOL. 10:293-300(1991).
CC -1- TISSUE SPECIFICITY: TESTIS, SPLEEN, KIDNEY, LUNG, HEART, AND BRAIN
CC (LOWEST LEVEL IN TESTIS AND HIGHEST IN LUNG).
CC -1- INDUCTION: BY GROWTH FACTORS.
CC -1- SIMILARITY: BELONGS TO THE INSULIN-LIKE GROWTH FACTOR BINDING
CC PROTEIN FAMILY. CEF-10/CYR61/CTGF/FISP-12/NOV PROTEIN SUBFAMILY.
CC -1- SIMILARITY: CONTAINS 1 VWFC DOMAIN.
CC -1- SIMILARITY: CONTAINS 1 C-TERMINAL CYSTINE KNOT-LIKE DOMAIN (CTCK).
DR EMBL; M70641; G193314; -.
DR EMBL; M70642; G193316; -.
DR EMBL; M80263; G201946; -.
DR PIR; A53228; A53228.
DR MGD; MGI:95537; FISP12.
DR PROSITE; PS00222; IGF-BINDING; 1.
DR PROSITE; PS01185; CTCK_1; 1.
DR PROSITE; PS01225; CTCK_2; 1.
DR PROSITE; PS01208; VWFC; 1.
KW GROWTH FACTOR BINDING; SIGNAL.
FT SIGNAL 1 25 POTENTIAL.
FT CHAIN 26 348 CONNECTIVE TISSUE GROWTH FACTOR.
FT DOMAIN 100 166 VWFC.
FT DOMAIN 255 329 CTCK.
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FT DISULFID 255 292 BY SIMILARITY.
FT DISULFID 272 306 BY SIMILARITY.
FT DISULFID 283 322 BY SIMILARITY.
FT DISULFID 286 324 BY SIMILARITY.
FT DISULFID 291 328 BY SIMILARITY.
FT CONFLICT 161 161 K -> E (IN REF. 2).
SQ SEQUENCE 348 AA; 37793 MW; EAB92BEO CRC32;

Query Match 12.2%; Score 112; DB 1; Length 348;
Best Local Similarity 30.4%; Pred. No. 4.8e-05;
Matches 34; Conservative 15; Mismatches 35; Indels 28; Gaps 7;

QY 27 DCPHQHDSSECKSSPRCK---RVLDGCGCRVCAAGRTCYRTVSGDMGKCGP--GL 81
DB 27 DCSAQCOCAA-EAAPHCPAGVSLVDGCGCRVCAKQLGELC-----TERDPCDPKHL 79
QY 82 RCPSNGEDPFGEFGICKD-----CPYG-----TFGMDCRETCNCSG 120
DB 80 FDFGS---PANRKIGVCTAKDGAFCVFGSVYRSGESFQSSCKYQCTCLDG 128

RESULT 8
IBP5_RAT
ID IBP5_RAT STANDARD; PRT; 271 AA.
AC P24594;
DT 01-MAR-1992 (REL. 21, CREATED)
DT 01-MAR-1992 (REL. 21, LAST SEQUENCE UPDATE)
DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)
DE INSULIN-LIKE GROWTH FACTOR BINDING PROTEIN 5 PRECURSOR (IGFBP-5)
DE (IBP-5) (IGF-BINDING PROTEIN 5).
GN IGFBP5 OR IGFBP-5.
OS RATTUS NORVEGICUS (RAT).
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;
OC EUTHERIA; RODENTIA.
RN [1]
RP SEQUENCE FROM N.A., AND SEQUENCE OF 20-53.
RX TISSUE-OVARY;
RX MEDLINE; 91244847.
RA SHIMASAKI S., SHIMONAKA M., ZHANG H.-P., LING N.;
RL J. BIOL. CHEM. 266:10646-10653(1991).
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN-SPRAGUE-DAWLEY;
RX MEDLINE; 93176146.
RA ZHU X., LING N., SHIMASAKI S.;
RL BIOCHEM. BIOPHYS. RES. COMMUN. 190:1045-1052(1993).
CC -1- FUNCTION: IGF-BINDING PROTEINS PROLONG THE HALF-LIFE OF THE IGFS
CC AND HAVE BEEN SHOWN TO EITHER INHIBIT OR STIMULATE THE GROWTH
CC PROMOTING EFFECTS OF THE IGFS ON CELL CULTURE. THEY ALTER THE
CC INTERACTION OF IGFS WITH THEIR CELL SURFACE RECEPTORS.
CC -1- TISSUE SPECIFICITY: MOSTLY IN KIDNEY.
CC -1- SIMILARITY: BELONGS TO THE INSULIN-LIKE GROWTH FACTOR BINDING
CC PROTEIN FAMILY.
DR EMBL; M62781; G204746; -.
DR EMBL; L08275; E73333; -.
DR PIR; A40403; A40403.
DR PIR; JCI463; JCI463.
DR PIR; F40403; F40403.
DR PROSITE; PS00222; IGF-BINDING; 1.
DR PROSITE; PS00484; THYROGLOBULIN_1; 1.
KW GROWTH FACTOR BINDING; SIGNAL.
FT SIGNAL 1 19 POTENTIAL.
FT CHAIN 20 271 INSULIN-LIKE GROWTH FACTOR BINDING
FT DOMAIN 214 262 THYROGLOBULIN TYPE 1.
SQ SEQUENCE 271 AA; 30298 MW; OAA79506 CRC32;

Query Match 12.0%; Score 110.5; DB 1; Length 271;
Best Local Similarity 28.7%; Pred. No. 5.8e-05;
Matches 27; Conservative 20; Mismatches 26; Indels 21; Gaps 3;
```

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01-OCT-1996 (REL. 34, LAST ANNOTATION UPDATE)
INSULIN-LIKE GROWTH FACTOR BINDING PROTEIN 5 PRECURSOR (IGFBP-5)
(IBF-5) (IGF-BINDING PROTEIN 5).
IGFBP5 OR IBP5.
HOMO SAPIENS (HUMAN).
EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;
EUETHERIA; PRIMATES.
[1]
SEQUENCE FROM N.A.
TISSUE=OSTEOSARCOMA;
RC MEDLINE: 91207396
RA KIEFER M.C., IOH R.S., BAUER D.M., ZAPF J.;
RL BIOCHEM. BIOPHYS. RES. COMMUN. 176:219-225(1991).
RN [2]
SEQUENCE FROM N.A.
TISSUE=PLACENTA;
RC MEDLINE: 91244847
RA SHIMASAKI S., SHIMOYAMA M., ZHANG H.-P., LING N.;
RL J. BIOL. CHEM. 266:10646-10653(1991).
RN [3]
SEQUENCE FROM N.A.
MEDLINE: 94193798
RA ALLANDER S.V., LARSSON C., EHRENBORG E., SUWANICHKUL A., WEBER G.,
RR MORRIS S.L., BAJALICK S., KIEFER M.C., LUTTMAN H., POWELL D.R.;
RL J. BIOL. CHEM. 269:10891-10898(1994).
RN [4]
SEQUENCE OF 24-43.
RX MEDLINE: 91207395.
RA ANDRESS D.L., BIRNBAUM R.S.;
RL BIOCHEM. BIOPHYS. RES. COMMUN. 176:213-218(1991).
CC -!- FUNCTION: IGF-BINDING PROTEINS PROLONG THE HALF-LIFE OF THE IGFS
AND HAVE BEEN SHOWN TO EITHER INHIBIT OR STIMULATE THE GROWTH
PROMOTING EFFECTS OF THE IGFS ON CELL CULTURE. THEY ALTER THE
INTERACTION OF IGFS WITH THEIR CELL SURFACE RECEPTORS.
CC -!- TISSUE SPECIFICITY: OSTEOSARCOMA, AND AT LOWER LEVELS IN LIVER,
KIDNEY, AND BRAIN.
CC -!- SIMILARITY: BELONGS TO THE INSULIN-LIKE GROWTH FACTOR BINDING
PROTEIN FAMILY.
DR EMBL: M65062; GI84820; -
DR EMBL: M62782; GI84818; -
DR EMBL: L27559; G505589; -
DR EMBL: L27556; G505589; JOINED.
DR EMBL: L27557; G505589; JOINED.
DR EMBL: L27558; G505589; JOINED.
DR PIR: JH0391; JH0391.
DR PIR: B40403; B40403.
DR PIR: PH0143; PH0143.
DR MIN: 146734; -.
DR PROSITE: PS00222; IGF_BINDING_1.
DR PROSITE: PS00484; THYROGLOBULIN_L1; 1.
KW GROWTH FACTOR BINDING; SIGNAL; POTENTIAL.
FT SIGNAL 1 20
CHAIN 21 272
INSULIN-LIKE GROWTH FACTOR BINDING
DOMAIN 215 263
SEQUENCE 272 AA; 30570 MW; 3AGAEF2 CRC32;
THYROGLOBULIN TYPE I.
PROTEIN 5.

Query Match 11.9%; Score 109.5; DB 1; Length 272;
Best Local Similarity 27.7%; Pred. No. 7.4e-05;
Matches 26; Conservative 20; Mismatches 27; Indels 21; Gaps

QY 4 VLLLTTLVPAHLVAWSNNYAVDCPHQCDSSECKSPKRRTVLDD-----DQGCRV 56
      :|||:: ||: :: |:::: |: :|::|
Db   8 LLALLAYAGPAQLSGSFV-----HCPCDEKALSMCPPSLGCCLVKPGCGCCMT 58
      :||||: ||: :: |:::: |: :|::|

QY 57 CAAGRGETCYRTVSMDGMKCPGLRCQPSNGED 90
      ||::: ||: :: |:::: |: :|::|
Db   59 CALAEGSC-----GVYTTERCAOGLCLRQDEE 87
      ||::: ||: :: |:::: |: :|::|

RESULT 11
CYR6 HUMAN
```

QY 106 ---TEGMDCRETCNCSG 120
Db 109 NGEFQPNCKHOCTCIDG 126

RESULT 12
CTGF_HUMAN
ID CTGF_HUMAN STANDARD; PRT; 349 AA.
AC P29279;
DT 01-DEC-1992 (REL. 24, CREATED)
DT 01-DEC-1992 (REL. 24, LAST SEQUENCE UPDATE)
DT 15-JUL-1998 (REL. 36, LAST ANNOTATION UPDATE)
DE CONNECTIVE TISSUE GROWTH FACTOR PRECURSOR.
GN CTGF.
OS HOMO SAPIENS (HUMAN).
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;
OC EUTHERIA; PRIMATES.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-UMBILICAL VEIN ENDOTHELIAL CELLS;
RX BRADHAM D.M., IGARASHI A., POTTER R.L., GROTEENDORST G.R.;
RL J. CELL BIOL. 114:1285-1294(1991).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE-UMBILICAL VEIN ENDOTHELIAL CELLS;
RX MEDLINE; 93187114.
RA IGARASHI A., BRADHAM D.M., OKOCHI H., GROTEENDORST G.R.;
RL J. DERMATOL. 19:642-643(1992).
RN [3]
RP SEQUENCE FROM N.A.
RA OSMAR B.S., WERNER A., YANG Z., GARNIER J.M., GENTZ R., LUESCHER T.F.;
RL SUBMITTED (APR-1994) TO EMBL/GENBANK/DBJ DATA BANKS.
CC -!- FUNCTION: MAJOR CONNECTIVE TISSUE MITOATTRACTANT SECRETED BY
HUMAN VASCULAR ENDOTHELIAL CELLS. THIS IMMEDIATE-EARLY PROTEIN
MAY BIND ONE OF THE PDGF CELL SURFACE RECEPTORS.
CC -!- SUBUNIT: MONOMER.
CC -!- ALTERNATIVE PRODUCTS: A SHORTER FORM MAY BE PRODUCED BY
ALTERNATIVE SPLICING OF THE SAME GENE.
CC -!- SIMILARITY: BELONGS TO THE INSULIN-LIKE GROWTH FACTOR BINDING
PROTEIN FAMILY. CEF-10/CYR61/CTGF/FTSP-12/NOV PROTEIN SUBFAMILY.
CC -!- SIMILARITY: CONTAINS 1 VFNC DOMAIN.
CC -!- SIMILARITY: CONTAINS 1 C-TERMINAL CYSTINE KNOT-LIKE DOMAIN (CTCK).
DR EMBL; M92934; G180924; -.
DR EMBL; X78947; G474934; -.
DR PIR; A40551; A40551.
DR PIR; S44205; S44205.
DR MIM; 121009; -.
DR PROSITE; PS00222; IGF_BINDING; 1.
DR PROSITE; PS01185; CTCK_1; 1.
DR PROSITE; PS01225; CTCK_2; 1.
DR PROSITE; PS01208; VFNC; 1.
DR KW GROWTH FACTOR BINDING; SIGNAL; ALTERNATIVE SPLICING.
FT SIGNAL 1 26
FT CHAIN 27 349
FT DOMAIN 101 167
FT DOMAIN 256 330
FT DISULFID 256 293
FT DISULFID 273 307
FT DISULFID 284 323
FT DISULFID 287 325
FT DISULFID 292 329
FT CARBOHYD 28 28
FT CARBOHYD 225 225
FT VARSPLIC 172 198
SQ SEQUENCE 349 AA; 38069 MW; C21E9662 CRC32;

Query Match 12.0%; Score 110.5; DB 1; Length 381;
Best Local Similarity 26.1%; Pred. No. 7.4e-05;
Matches 36; Conservative 21; Mismatches 44; Indels 37; Gaps 8;

QY 2 KSVLLTLLVPAHLVAWSNNYAVDCPOHCDSSCKSPCKRT---VLDGCGCRVCA 58
Db 7 RALALVVTLLHLTRLALS-----TCPAACHCP-LEAPKCAPGVGLRDGCGCKVCA 57

QY 59 AGRGETCYRTVSGMDGKCGP--GLRCQPSNGEDPFGEFGICKD-----CPYG----- 106
Db 58 KOLNEDCSKTOP-----CDHTKGLCNEFGASSTALK---GICRAQSEGRPCFCEYNSRIYQ 108

Query Match 11.9%; Score 109; DB 1; Length 349;
Best Local Similarity 30.4%; Pred. No. 0.0001;
Matches 34; Conservative 14; Mismatches 36; Indels 28; Gaps 7;

Db 28 NCSGPCRCD-EPAPRCAGVSLVLDGCGCRVCAKOLGELC-----TERDFCDPHKGL 80
 QY 82 RCQPSNGEDPFGEEFGCKD-----CPYG-----TFGMDCRETNCOSQ 120
 Db 81 FCDFGS---PANKIGVCTAKDAGPCIFGTVYRSGESFQSSCKYQCTCLDG 129

RESULT 13
 IBP4_SHEEP
 ID IBP4_SHEEP STANDARD; PRT; 237 AA.
 AC Q28893;
 DT 01-NOV-1997 (REL. 35, CREATED)
 DT 01-NOV-1997 (REL. 35, LAST SEQUENCE UPDATE)
 DE 01-NOV-1997 (REL. 35, LAST ANNOTATION UPDATE)
 DE INSULIN-LIKE GROWTH FACTOR BINDING PROTEIN 4 (IGFBP-4) (IBP-4)
 GN (IGF-BINDING PROTEIN 4).
 OS IGFBP4.
 OS OVIS ARIES (SHEEP).
 OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;
 OC EUTHERIA; ARTIODACTYLA.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE-LIVER;
 RX MEDLINE; 95151165.
 RA CARR J.M., GRANT P.A., FRANCIS G.L., OWENS J.A., WALLACE J.C.,
 RA WALTON P.E.;
 RL J. MOL. ENDOCRINOL. 13:219-236(1994).

CC -1- FUNCTION: IGF-BINDING PROTEINS PROLONG THE HALF-LIFE OF THE IGFS
 AND HAVE BEEN SHOWN TO EITHER INHIBIT OR STIMULATE THE GROWTH
 CC PROMOTING EFFECTS OF THE IGFS ON CELL CULTURE. THEY ALTER THE
 CC INTERACTION OF IGFS WITH THEIR CELL SURFACE RECEPTORS.
 CC -1- SIMILARITY: BELONGS TO THE INSULIN-LIKE GROWTH FACTOR BINDING
 CC PROTEIN FAMILY.
 DR EMBL; S77394; G944952; -;
 DR PROSITE; PS00222; IGF_BINDING; 1.
 DR PROSITE; PS00484; THYROGLOBULIN_1; 1.
 KW GROWTH FACTOR BINDING; GLYCOPROTEIN.
 FT DOMAIN 179 228 THYROGLOBULIN TYPE I.
 FT CARBOHYD 104 104 POTENTIAL.
 SQ SEQUENCE 237 AA; 25869 MW; C1C79FEA CRC32;

Query Match 11.7%; Score 107; DB 1; Length 237;
 Best Local Similarity 34.8%; Pred. No. 0.00012;
 Matches 24; Conservative 13; Mismatches 24; Indels 8; Gaps 3;

QY 25 AVDCPQHCHDS--SECKSPRCRTVLD-DCGCRVCAAGRETCTYVSGMDGMKCGPGL 81
 Db 3 AIHCPPESEKLCARCPVGCCELVRPGCGCCATCALGKGMPC-----GVYTPDCGSGL 57
 QY 82 RCQPSNGED 90
 Db 58 RCHPFRGVE 66

RESULT 14
 IBP4_HUMAN
 ID IBP4_HUMAN STANDARD; PRT; 258 AA.
 AC P22692;
 DT 01-AUG-1991 (REL. 19, CREATED)
 DT 01-MAR-1992 (REL. 21, LAST SEQUENCE UPDATE)
 DE 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)
 DE INSULIN-LIKE GROWTH FACTOR BINDING PROTEIN 4 (IGFBP-4)
 GN (IBP-4) (IGF-BINDING PROTEIN 4).
 GN IGFBP4 OR IBP4.
 OS HOMO SAPIENS (HUMAN).
 OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;
 OC EUTHERIA; PRIMATES.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 91186988.
 RA LATOUR D., MOHAN S., LINKHART T.A., BAYLINK D.J., STRONG D.D.;

RL MOL. ENDOCRINOL. 4:1806-1814(1990).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE-PLACENTA;
 RX MEDLINE; 91133415.
 RA SHIMASAKI S., UCHIYAMA F., SHIMONAKA M., LING N.;
 RL MOL. ENDOCRINOL. 4:1451-1458(1990).
 RN [3]
 RP SEQUENCE FROM N.A., AND SEQUENCE OF 22-41.
 RC TISSUE-OSTEOSARCOMA;
 RX MEDLINE; 91225006.
 RA KIEFER M.C., MASIARZ F.R., BAUER D.M., ZAPF J.;
 RL J. BIOL. CHEM. 266:9043-9049(1991).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC TISSUE-PLACENTA;
 RA STRONG D.D., MORALES S., LEE K., BOONYARATANAKORNKIT V.,
 RA BAYLINK D.J., MOHAN S.;
 RL SUBMITTED (FEB-1995) TO EMBL/GENBANK/DBJ DATA BANKS.
 RN [5]
 RP SEQUENCE OF 22-53.
 RC TISSUE-COLON;
 RX MEDLINE; 91235178.
 RA CULOUSCOU J.-M., SHOYAB M.;
 RL CANCER RES. 51:2813-2819(1991).
 CC -1- FUNCTION: IGF-BINDING PROTEINS PROLONG THE HALF-LIFE OF THE IGFS
 AND HAVE BEEN SHOWN TO EITHER INHIBIT OR STIMULATE THE GROWTH
 CC PROMOTING EFFECTS OF THE IGFS ON CELL CULTURE. THEY ALTER THE
 CC INTERACTION OF IGFS WITH THEIR CELL SURFACE RECEPTORS.
 CC -1- BINDS IGF-II MORE THAN IGF-I.
 CC -1- SIMILARITY: BELONGS TO THE INSULIN-LIKE GROWTH FACTOR BINDING
 CC PROTEIN FAMILY.
 DR EMBL; M38177; -; NOT_ANNOTATED_CDS.
 DR EMBL; M62403; G184816; -;
 DR EMBL; U20982; G695254; -;
 DR PIR; A36549; A36549.
 DR PIR; B37252; B37252.
 DR PIR; B39842; B39842.
 DR MIM; 146733; -;
 DR PROSITE; PS00222; IGF_BINDING; 1.
 DR PROSITE; PS00484; THYROGLOBULIN_1; 1.
 KW GROWTH FACTOR BINDING; SIGNAL; GLYCOPROTEIN.
 FT SIGNAL 1 21
 FT CHAIN 22 258 INSULIN-LIKE GROWTH FACTOR BINDING
 FT CARBOHYD 125 125 POTENTIAL.
 FT DOMAIN 200 249 THYROGLOBULIN TYPE I.
 FT CONFLICT 51 51
 FT CONFLICT 198 198 P -> A (IN REF. 1, 4 AND 5).
 FT CONFLICT 198 198 I -> F (IN REF. 1 AND 4).
 SQ SEQUENCE 258 AA; 27934 MW; 58AC8AC3 CRC32;

Query Match 11.6%; Score 106.5; DB 1; Length 258;
 Best Local Similarity 31.8%; Pred. No. 0.00015;
 Matches 28; Conservative 22; Mismatches 29; Indels 9; Gaps 4;

QY 6 LLTLLVPAHLVAWSNNYAVDCPQHCHDS--SECKSPRCRTVLD-DCGCRVCAAGRG 62
 Db 6 LVAALLAAGPGLSLGDE-AIHCPPESEKLCARCPVGCCELVRPGCGCCATCALGLG 64
 QY 63 ETCYRTVSGMDGMKCGPGLRCQPSNGED 90
 Db 65 MPC-----GVYTPRCGSLRCYPPRGVE 87

RESULT 15
 IBP3_PIG
 ID IBP3_PIG STANDARD; PRT; 266 AA.
 AC P16611;
 DT 01-AUG-1990 (REL. 15, CREATED)
 DT 01-FEB-1991 (REL. 17, LAST SEQUENCE UPDATE)
 DT 01-FEB-1996 (REL. 33, LAST ANNOTATION UPDATE)
 DE INSULIN-LIKE GROWTH FACTOR BINDING PROTEIN 3 (IGFBP-3) (IBP-3) (IGF-

DE BINDING PROTEIN 3).
GN IGFBP3.
OS SUS SCROFA (PIG).
OC EURARYOTA; METAZOA; CHORDATA; VERTEBRATA; TETRAPODA; MAMMALIA;
CC EUTHERIA; ARTIODACTYLA.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 90130475.
RA SHIMASAKI S., SHIMONAKA M., UI M., INOUE S., SHIBATA F., LING N.;
RL J. BIOL. CHEM. 265:2198-2202(1990).
RN [2]
RP SEQUENCE OF 1-15.
RX MEDLINE: 92109718.
RA COLEMAN M.E., PAN Y.-C.E., ETHERTON T.D.;
RL BIOCHEM. BIOPHYS. RES. COMMUN. 181:1131-1136(1991).
CC -1- FUNCTION: IGF-BINDING PROTEINS PROLONG THE HALF-LIFE OF THE IGFS
CC AND HAVE BEEN SHOWN TO EITHER INHIBIT OR STIMULATE THE GROWTH
CC PROMOTING EFFECTS OF THE IGFS ON CELL CULTURE. THEY ALTER THE
CC INTERACTION OF IGFS WITH THEIR CELL SURFACE RECEPTORS.
CC -1- SUBUNIT: FORMS A TERNARY COMPLEX OF ABOUT 140 TO 150 KD WITH IGF-I
CC OR IGF-II AND A 85 KD GLYCOPROTEIN (ALS).
CC -1- SUBCELLULAR LOCATION: EXTRACELLULAR.
CC -1- BINDS IGF-II MORE THAN IGF-I.
CC -1- SIMILARITY: BELONGS TO THE INSULIN-LIKE GROWTH FACTOR BINDING
CC PROTEIN FAMILY.
CC EMBL: J05228; G164501; .
DR PIR: A35037; A35037.
DR PIR: JH0516; JH0516.
DR HSP; P17494; IKST.
DR PROSITE; PS00222; IGF_BINDING; 1.
DR PROSITE; PS00484; THYROGLOBULIN.1; 1.
KW GROWTH FACTOR BINDING; GLYCOPROTEIN.
FT CARBOHYD 91 91 POTENTIAL.
FT CARBOHYD 111 111 POTENTIAL.
FT CARBOHYD 174 174 POTENTIAL.
FT DOMAIN 211 260 THYROGLOBULIN TYPE I.
FT CONFLICT 5 6 VG -> A (IN REF. 2).
SQ SEQUENCE 266 AA; 28910 MW; 53E9830D CRC32;

Query Match 11.4%; Score 104.5; DB 1; Length 266;
Best Local Similarity 31.3%; Pred. No. 0.00025;
Matches 26; Conservative 18; Mismatches 20; Indels 19; Gaps 4;
QY 31 HCDSSCK-----SSPRCKRTVLD-DCGCRVCAAGRCETCYRTVSGMDGMCQPG 80
Db 12 RCEPCDARALAQAPPAAPCAELVREPCGCGCLTALREGQAC-----GVYTERCGAG 66
QY 81 LRCQPSNGE----DPFGEEFGIC 99
Db 67 LRCQPPPGEPRLQALLDGRGIC 89

Search completed: May 4, 1999, 16:50:02
Job time: 10240 sec

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OM protein - protein search, using sw model

Run on: May 4, 1999, 02:47:21 ; Search time 36.71 Seconds
(without alignments)
276.522 Million cell updates/sec

Title: US-09-037-460-2

Perfect score: 184

Sequence: 1 MKSVLLLTLLVPAHLVAWA.....EYVKENAGSPVWRKWLNPR 184

Scoring table: OLIGO

Searched: 180763 seqs, 55169189 residues

Database: SPTRMBL8:

- 1: sp_fungi:*
- 2: sp_human:*
- 3: sp_invertebrate:*
- 4: sp_mammal:*
- 5: sp_mhc:*
- 6: sp_organelle:*
- 7: sp_phage:*
- 8: sp_plant:*
- 9: sp_bacteria:*
- 10: sp_rodent:*
- 11: sp_virus:*
- 12: sp_vertebrate:*
- 13: sp_unclassified:*
- 14: sp_archaea:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	184	100.0	184	2 Q15330	Q15330 homo sapien

ALIGNMENTS

RESULT 1
Q15330
ID Q15330 PRELIMINARY; PRT; 184 AA.
AC Q15330;
DT 01-NOV-1996 (TREMBLREL. 01, CREATED)
DT 01-NOV-1996 (TREMBLREL. 01, LAST SEQUENCE UPDATE)
DT 01-NOV-1998 (TREMBLREL. 08, LAST ANNOTATION UPDATE)
DE ESM-1 SECRETORY PROTEIN PRECURSOR.
OS HOMO SAPIENS (HUMAN).
OC EUKARYOTA; METAZOA; CHORDATA; VERTEBRATA; MAMMALIA; EUTHERIA; PRIMATES;
CC CATARRHINI; HOMINIDAE; HOMO.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 96355375.
RA LASSALLE P.M., MOLET S., JANIN A., VANDER-HEYDEN J.E., TAVERNIER J.,
RT FIERIS W., DEVOS R.E., TONNEL A.E.B.;
RT "ESM-1 is a novel human endothelial cell-specific molecule expressed
in lung and regulated by cytokines";
RL J. BIOL. CHEM. 271:20458-20464(1996).
DR EMBL; X89426; E189266; -;
DR PFAM; PF00219; IGFBP; 1.
KW SIGNAL.
FT SIGNAL 1 19 POTENTIAL.

FT CHAIN 20 184 ESM-1 SECRETORY PROTEIN.
SQ SEQUENCE 184 AA; 20095 MW; 08D109DF CRC32;

Query Match 100.0%; Score 184; DB 2; Length 184;
Best Local Similarity 100.0%; Pred. No. 1.5e-180;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MKSVLLLTLLVPAHLVAWNNYAVDCPQHCDSSECKSSPRCKRTVLDGCGCRVCAAG 60
Db 1 MKSVLLLTLLVPAHLVAWNNYAVDCPQHCDSSECKSSPRCKRTVLDGCGCRVCAAG 60
QY 61 RGETCYRTVSGMDGKCGPLRCOPSGNEDPFGEEFGICKDCPYGTGMDCRETNCQSG 120
Db 61 RGETCYRTVSGMDGKCGPLRCOPSGNEDPFGEEFGICKDCPYGTGMDCRETNCQSG 120
QY 121 ICDRGTKCLKFFPFQYSVTKSSNRFVSLTEHDMASGDGNIVREEVYKNAAGSPVWRKW 180
Db 121 ICDRGTKCLKFFPFQYSVTKSSNRFVSLTEHDMASGDGNIVREEVYKNAAGSPVWRKW 180
QY 181 LNPR 184
Db 181 LNPR 184

Search completed: May 4, 1999, 05:33:53
Job time: 9992 sec

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OM protein - protein search, using sw model

Run on: May 4, 1999, 09:00:24 ; Search time 26.81 Seconds
(without alignments)

Title: US-09-037-460-2

Perfect score: 184

Sequence: 1 MKSVLLLTLLVPAHLVAW.....EVVKENAAGSPVMRKWLNPR 184

Scoring table: OLIGO

Searched: 162890 seqs, 20225328 residues

Database : A_Geneseq_34:**

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

word = 30

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	184	100.0	184	1 R98994	Vascular IBP-like

ALIGNMENTS

RESULT 1

R98994	R98994 standard; Protein; 184 AA.
ID	AC R98994;
DE	06-NOV-1996 (first entry)
DT	Vascular IBP-like growth factor.
DD	Vascular IBP-like growth factor; VIGF;
DE	insulin-like growth factor binding protein; agonist; antagonist;
KW	muscle wastage; osteoporosis; implant fixation; wound healing;
KW	therapy; diagnosis.
KW	Homo sapiens.
OS	Homo sapiens.
EH	Key
FT	Location/Qualifiers
FT	1..21
FT	/label= Sig_Peptide
PN	W09617931-AL.
PD	13-JUN-1996.
PP	09-DEC-1994; U14388.
PR	09-DEC-1994; WO-U14388.
PP	(HUMA-) HUMAN GENOME SCI INC.
PI	Hastings GA, Rosen CA;
PI	WPI; 96-287176/29.
PI	N-PSDB; T34991.
DD	Human vascular insulin-like growth factor binding protein-like
PPT	growth factor, and its nucleic acid sequence and (ant)agonists
PPT	used, e.g. to treat muscle wasting diseases or aid implant fixation,
PPT	or limit excess connective tissue prodn. during wound healing.
PPS	Claim 14; Page 43-44; 61pp; English.
CCC	Human vascular insulin-like growth factor binding protein-like
CCC	growth factor [R98994], or VIGF, is a protein of primarily
CCC	vascular origin that is structurally related to the IBP and CCN
CCC	protein families. It can be expressed in e.g. E. coli, CHO or
CCC	insect host cells using a vector incorporating a cDNA clone
CCC	[R34991], or its derivative, obtd. from human umbilical
CCC	endothelial cells. It is useful therapeutically e.g. for
CCC	treating muscle wasting diseases or osteoporosis, or can be used
CCC	to detect diseases associated with under- or over-expression of VIGF,
CCC	or to screen for antagonists useful during wound healing.

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OM protein - protein search, using sw model

Run on: May 4, 1999, 08:18:16 ; Search time 25.07 Seconds
(without alignments)

Title: US-09-037-460-2_COPY_55_69
Perfect score: 78
Sequence: 1 RVCAAGRGETCYRTV 15

Scoring table: PAM150

Searched: 116738 seqs, 37460341 residues

```
Database : PIR_58:*
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*
```

SUMMARIES

Result No.	Score	Query %		DB	ID	Description
		Match	Length			
1	40	51.3	115	2	I37901	Histone H2A relate
2	42	53.8	607	1	ABHOS	serum albumin prec
3	39	50.0	218	2	A70080	two-component resp
4	36	46.2	107	1	A60361	neuroparsin A prec
5	36	46.2	136	2	S78428	destabilase 2 - me
6	31	39.7	25	2	JH0701	omega-conotoxin MV
7	29	37.2	12	2	S26553	T-cell receptor be
8	33.5	42.9	81	2	JC4621	cardiotoxin N prec
9	38	48.7	599	2	A36100	genome polyprotein
10	29.5	37.8	17	2	A38824	tachyplesin III -
11	29.5	37.8	17	2	JX0125	tachyplesin III -
12	29.5	37.8	17	2	A30068	tachyplesin II - h
13	29.5	37.8	17	2	JU0123	tachyplesin II - h
14	35	44.9	185	2	S59504	ferric pseudobacti
15	29.5	37.8	19	2	JX0134	tachyplesin I prec
16	39	50.0	1040	2	E71412	hypothetical prote
17	34	43.6	130	2	S52116	transforming prote
18	38	48.7	728	1	JH0579	hepatocyte growth
19	33.5	42.9	122	2	A25806	phospholipase A2 (
20	36	46.2	351	2	S20078	NOV protein - chic
21	36	46.2	357	2	I38069	gene novH protein
22	28	35.9	13	2	B58533	CD61 homolog - cha
23	36	46.2	387	3	JC5949	galanin receptor 2
24	36	46.2	388	2	C70605	probable aminotran
25	33.5	42.9	138	1	PSV1AA	phospholipase A2 (
26	33.5	42.9	138	1	PSV1AA	phospholipase A2 (
27	33.5	42.9	138	2	S10333	ammodytoxin B prec
28	33.5	42.9	138	2	I51386	phospholipase A2 (
29	35	46.2	395	2	S40685	probable G protein
30	31.5	40.4	60	1	H3NJ21	cytotoxin 2 - Naja
31	31.5	40.4	60	1	H3NJ2R	cytotoxin 2 - Asia
32	31.5	40.4	60	1	H3NJ2F	cytotoxin 2 - Chin
33	31.5	40.4	60	1	H3NJ4F	cytotoxin 4 - Chin
34	31.5	40.4	60	1	H3NJ5F	cytotoxin 5 - Chin
35	31.5	40.4	60	1	H3NJ2K	cytotoxin 2 - mono
36	31.5	40.4	60	1	H3NJ3K	cytotoxin 3 - mono
37	31.5	40.4	60	1	H3NJ1I	cytotoxin 1 - Naja
38	31.5	40.4	60	1	H3NJ1K	cytotoxin 1 - mono
39	31.5	40.4	60	2	JS0028	cytotoxin II - mon

ALIGNMENTS

RESULT 1
I37901
Histone H2A related - human
C:Species: Homo sapiens (man)
C:Date: 29-May-1998 #sequence_revision 29-May-1998 #text_change 21-Aug-1998
C:Accession: I37901
F:Naylor, J.A.; Buck, D.; Green, P.M.; Williamson, H.; Bentley, D.; Giannelli, F.
Hum. Mol. Genet. 4, 1217-1224, 1995
A:Title: Investigation of the Factor VIII intron 22 repeated region (int22h) and the
A:Reference number: I37901
A:Accession: I37901
A>Status: preliminary; translated from GB/EMBL/DDBJ
A:Molecule type: DNA
A:Residues: 1-115 <RES>
A:Cross-references: EMBL:X86012; NID:g976082; PID:g976083
C:Superfamily: histone H2A

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QY      4  AAGRGETCYRTV 15
.      : ||| || |||
DB     13  AGGRGRTCSRTV 24

RESULT 2
ABHOS
serum albumin precursor - horse
C:Species: Equus caballus (domestic horse)
C:Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 05-Sep-1997
C:Accession: S34053
R:Ho, J.X.; Holowachuk, E.W.; Norton, E.J.; Twigg, P.D.; Carter, D.C.
Eur. J. Biochem. 215, 205-212, 1993
A:Title: X-ray and primary structure of horse serum albumin (Equus caballus) at 0.27-
A:Reference number: S34053
A:Accession: S34053
A:Molecule type: mRNA
A:Residues: 1-607 <HOA>
A:Cross-references: GB:X74045; NID:G399671; PID:G399672
C:Comment: Serum albumin is synthesized in the liver as preproalbumin. It binds copper
teroid hormones (weak bonds with these hormones promote their transfer across the mem
C:Superfamily: serum albumin; serum albumin repeat homology
C:Keywords: carrier protein; duplication; metal binding; plasma
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-24/Domain: signal sequence #status predicted <PRO>
F:25-607/Product: serum albumin #status predicted <MAT>
F:29-201/Domain: serum albumin repeat homology <SA1>
F:220-393/Domain: serum albumin repeat homology <SA2>
F:412-591/Domain: serum albumin repeat homology <SA3>
F:27/Binding site: copper (His) #status predicted
F:77-86,99-115,114-125,147-192,191-200,223-269,288-276,288-302,301-312,339-384,383-39
F:263/Binding site: bilirubin (Lys) #status predicted

Query Match      53.8%; Score 42; DB 1; Length 607;
Best Local Similarity 46.7%; Pred. No. 8;
Matches 7; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY      1  RVCAAGRGETCYRTV 15
.      : ||| : |||||

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QY 1 RVCAAGRGETCYRTV 15
: || : : || ||

Db 382 KCAEADPPACRYTV 396

RESULT 3

A70080
two-component response regulator [YxjM] homolog yxjL - Bacillus subtilis
C:Species: Bacillus subtilis
C>Date: 05-Dec-1997 #sequence_revision 05-Dec-1997 #text_change 24-Sep-1998
C:Accession: A70080
R:Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Bertoldo, C.; Bron, S.; Brouillet, S.; Bruschi, C.V.; Caldwell, B.; Capuano, V.; Carter, N.M.; Chao, A.; Ehrlich, S.D.; Emerson, P.T.; Entian, K.D.; Errington, J.; Fabret, C.; Ferrari, E.; Nature 390, 249-256, 1997
A:Authors: Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Gallizzi, A.; Galleron, N.; Ghim, wood, C.R.; Henaut, A.; Hilbert, H.; Hoisappel, S.; Hosono, S.; Hullo, M.F.; Itaya, M.; Koningsstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardinois, S.; Lauber, A.; Authors: Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Maueel, C.; Medigu, K.; Ogiwara, A.; Oudega, B.; Park, S.H.; Parro, V.; Pohl, T.M.; Portetalle, D.; Porwol, Rivolta, C.; Rocha, E.; Roche, B.; Rose, M.; Sadaie, Y.; Sato, T.; Scanlon, E.; Schleic, A:Authors: Schroeter, R.; Scoffone, F.; Sekiguchi, J.; Sekowska, A.; Seror, S.J.; Serro amakoshi, A.; Tanaka, T.; Terpstra, P.; Tognoni, A.; Yasumoto, K.; Yata, K.; Yoshida, K.; Yoshikawa, A.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yata, K.; Yoshida, K.; Yoshikawa, A.; Authors: Zumbstein, E.; Yoshikawa, H.; Danchin, A.
A:Title: The complete genome sequence of the Gram-positive bacterium Bacillus subtilis.
A:Reference number: A69580; MUID:98044033
A:Accession: A70080
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-218 <N>
A:Cross-references: GB:299123; GB:AL009126; NID:g2636240; PID:el186390; PID:g2636426
A:Experimental source: strain 168
C:Genetics:
A:Gene: yxjL
C:Superfamily: regulatory protein comA; response regulator homology
F:8-119/Domain: response regulator homology <RRH>
F:58/Binding site: phosphate (Asp) (covalent) #status predicted

Query Match 50.0%; Score 39; DB 2; Length 218;

Best Local Similarity 66.7%; Pred. No. 9.9;

Matches 8; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 4 AAGRGTCYRTV 15

II:III:IIII

Db 121 AARGAIFRTV 132

RESULT 4

A60361
neuroparsin A precursor - migratory locust
N:Contains: neuroparsin A; neuroparsin B
C:Species: Locusta migratoria (migratory locust)
C>Date: 30-Jun-1993 #sequence_revision 12-Apr-1996 #text_change 12-Apr-1996
C:Accession: A61618; A60361; S03545
R:Laqueux, M.; Kromer, E.; Girardie, J.
Insect Biochem. Mol. Biol. 22, 511-516, 1992
A:Title: Cloning of a Locusta cDNA encoding neuroparsin A.
A:Reference number: A61618
A:Accession: A61618
A:Status: not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-107 <LAG>
R:Girardie, J.; Huet, J.C.; Pernollet, J.C.
Insect Biochem. 20, 659-666, 1990
A:Title: The locust neuroparsin A: sequence and similarities with vertebrate and insect
A:Reference number: A60361
A:Accession: A60361
A:Molecule type: protein
A:Residues: 25-107 <GR>
R:Girardie, J.; Girardie, A.; Huet, J.C.; Pernollet, J.C.
FEBS Lett. 245, 4-8, 1989
A:Title: Amino acid sequence of locust neuroparsins.
A:Reference number: S03545; MUID:89171328

A:Accession: S03545

A:Molecule type: protein

A:Residues: 30-107 <GI2>

C:Comment: Intermediate forms of the neuroparsin monomer are also found. It is not kn

C:Superfamily: neuroparsin

C:Keywords: disulfide bond; homodimer; hormone

F:1-22/Domain: signal sequence #status predicted <SIG>

F:23-24/Domain: propeptide #status experimental <PRO>

F:25-107/Product: neuroparsin A monomer #status experimental <MAT1>

F:30-107/Product: neuroparsin B monomer #status experimental <MAT2>

Query Match 46.2%; Score 36; DB 1; Length 107;

Best Local Similarity 54.5%; Pred. No. 17;

Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 RVCAARGETC 11

:IIII:II

Db 55 KVCAGPGDKC 65

RESULT 5

S78428
destabilase 2 - medicinal leech
C:Species: Hirudo medicinalis (medicinal leech)
C>Date: 12-Feb-1998 #sequence_revision 13-Mar-1998 #text_change 13-Mar-1998
C:Accession: S78428
R:Frackov, A.; Berezhnoy, S.; Barsova, E.V.; Zavalova, L.; Lukyanov, S.; Baskova, I.; FEBS Lett. 390, 145-148, 1996
A:Title: Enzyme from the medicinal leech (Hirudo medicinalis) that specifically split
A:Reference number: S78427
A:Accession: S78428
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-136 <FRA>
A:Cross-references: EMBL:U24122; NID:gl255717; PID:gl255718

Query Match 46.2%; Score 36; DB 2; Length 136;

Best Local Similarity 54.5%; Pred. No. 21;

Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 RVCAARGETC 11

:IIII:II

Db 95 RFTGGRTPTC 105

RESULT 6

JH0701
omega-conotoxin MVIIB - cone shell (Conus magus)
C:Species: Conus magus (magus cone)
C>Date: 17-Apr-1993 #sequence_revision 17-Apr-1993 #text_change 23-May-1997
C:Accession: JH0701; B34115
R:Hillyard, D.R.; Monje, V.D.; Mintz, I.M.; Bean, B.P.; Nadasdi, L.; Ramachandran, J.
Neuron 9, 69-77, 1992
A:Title: A new conus peptide ligand for mammalian presynaptic Ca2+ channels.
A:Reference number: JH0699; MUID:92337922
A:Accession: JH0701
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-25 <HIL>
R:Olivera, B.M.; Cruz, L.J.; de Santos, V.; LeCheminant, G.W.; Griffin, D.; Zeikus, R.
Biochemistry 26, 2086-2090, 1987
A:Title: Neuronal calcium channel antagonists. Discrimination between calcium channel
A:Reference number: A34115; MUID:87299637
A:Accession: B34115
A:Molecule type: protein
A:Residues: 1-25 <OLI>
C:Superfamily: omega-conotoxin
C:Keywords: acetylcholine release inhibition; amidated carboxyl end; calcium channel
F:1-16,8-20,15-25/Disulfide bonds: #status predicted
F:25/Modified site: amidated carboxyl end (Cys) #status predicted

Query Match 39.7%; Score 31; DB 2; Length 25;
 Best Local Similarity 55.6%; Pred. No. 30;
 Matches 5; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 6 GRGETCYR 14
 Db 3 GKGASCHRT 11

RESULT 7
 T-cell receptor beta chain (clone Cw3/56.1) - mouse (fragment)
 C:Species: Mus musculus (house mouse)
 C:Date: 13-Jan-1995 #sequence_revision 17-Apr-1998 #text_change 17-Apr-1998
 C:Accession: S26553
 R:Casanova, J.L.; Cerottini, J.C.; Matthes, M.; Necker, A.; Gournier, H.; Barra, C.; Wid
 J. Exp. Med. 176, 439-447, 1992
 A:Title: H-2-restricted cytolytic T lymphocytes specific for HLA display T cell receptor
 A:Reference number: S26512
 A:Accession: S26553
 A:Molecule type: mRNA
 A:Residues: 1-12 <CAS>
 A:Cross-references: EMBL:X68003
 A:Experimental source: cytolytic T-lymphocyte, clone Cw3/56.1
 C:Superfamily: immunoglobulin V region; immunoglobulin homology
 C:Keywords: T-cell receptor

Query Match 37.2%; Score 29; DB 2; Length 12;
 Best Local Similarity 60.0%; Pred. No. 33;
 Matches 6; Conservative 3; Mismatches 3; Indels 1; Gaps 0;

OY 3 CAARGETCY 12
 Db 1 CASSWGTELY 10

RESULT 8
 JC4621
 cardiotoxin N precursor - Chinese cobra
 N:Alternate names: ctxn
 C:Species: Naja naja atra (Chinese cobra)
 C:Date: 10-Apr-1996 #sequence_revision 24-May-1996 #text_change 25-Apr-1997
 C:Accession: JC4621; JC2469
 R:Chang, L.S.; Wu, P.F.; Lin, J.
 Biochem. Biophys. Res. Commun. 219, 116-121, 1996
 A:Title: cDNA sequence analysis and expression of cardiotoxins from Taiwan cobra.
 A:Reference number: JC4619; MUID:96190679
 A:Accession: JC4621
 A:Molecule type: mRNA
 A:Residues: 1-81 <CHA>
 A:Cross-references: EMBL:Z54230; NID:g1054814; PID:g1000509
 A:Experimental source: venom glands
 R:Chiou, S.H.; Hung, C.C.; Huang, H.C.; Chen, S.T.; Wang, K.T.; Yang, C.C.
 Biochem. Biophys. Res. Commun. 206, 22-32, 1994
 A:Title: Sequence comparison and computer modelling of cardiotoxins and cobrotoxin iso
 A:Reference number: JC2469
 A:Accession: JC2469
 A:Molecule type: protein
 A:Residues: 22-81 <CHI>
 A:Note: conformation by (1)H-NMR
 C:Superfamily: snake toxin
 C:Keywords: cardiotoxin; venom
 F:1-21/Domain: signal sequence #status predicted <SIG>
 F:22-81/Product: cardiotoxin N #status experimental <WAT>
 F:24-42,35-59,63-74,75-80/Disulfide bonds: #status predicted

Query Match 42.9%; Score 33.5; DB 2; Length 81;
 Best Local Similarity 46.2%; Pred. No. 34;
 Matches 6; Conservative 4; Mismatches 2; Indels 1; Gaps 1;

OY 1 RVCAARGETCYR 13
 Db 33 KTCAGR-NLCYK 44

RESULT 9
 A26100
 genome polyprotein - murine poliovirus (fragment)
 N:Contains: proteinase (EC 3.4.-.-); RNA-directed RNA polymerase (EC 2.7.7.48)
 C:Species: murine poliovirus, Theiler's encephalomyelitis virus
 C:Date: 31-Mar-1991 #sequence_revision 31-Mar-1991 #text_change 05-Dec-1997
 C:Accession: A26100
 R:Ozden, S.; Taqay, F.; Chamorro, M.; Brahic, M.
 J. Virol. 60, 1163-1165, 1986
 A:Title: Theiler's virus genome is closely related to that of encephalomyocarditis vi
 A:Reference number: A26100; MUID:87061197
 A:Accession: A26100
 A:Molecule type: genomic RNA
 A:Residues: 1-599 <OZD>
 A:Cross-references: GB:M14703; NID:g335241; PID:g335242
 C:Superfamily: foot-and-mouth disease virus genome polyprotein
 C:Keywords: hydrolase; nucleotidyltransferase; polypeptide; proteinase
 F:1-139/Product: proteinase (fragment) #status predicted <PTS>
 F:140-599/Product: RNA-directed RNA polymerase #status predicted <RPS>

Query Match 48.7%; Score 38; DB 2; Length 599;
 Best Local Similarity 55.6%; Pred. No. 38;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

OY 5 AGRGETCYR 13
 Db 38 PARNDTCYR 46

RESULT 10
 A38824
 tachyplesin I - horseshoe crab (Tachyplesus gigas)
 C:Species: Tachyplesus gigas
 C:Date: 30-Jun-1992 #sequence_revision 30-Jun-1992 #text_change 11-Jul-1997
 C:Accession: A38824
 R:Muta, T.; Fujimoto, T.; Nakajima, H.; Iwanaga, S.
 J. Biochem. 108, 261-266, 1990
 A:Title: Tachyplesins isolated from hemocytes of southeast Asian horseshoe crabs (Car
 ssing intermediate of its precursor
 A:Reference number: JX0124; MUID:91035357
 A:Accession: A38824
 A:Molecule type: protein
 A:Residues: 1-17 <WUT>
 A:Experimental source: hemocyte
 C:Keywords: amidated carboxyl end
 F:3-16,7-12/Disulfide bonds: #status predicted
 F:17/Modified site: amidated carboxyl end (Arg) #status experimental

Query Match 37.8%; Score 29.5; DB 2; Length 17;
 Best Local Similarity 53.8%; Pred. No. 38;
 Matches 7; Conservative 0; Mismatches 3; Indels 3; Gaps 1;

OY 1 RVCAARGETCYR 13
 Db 5 RVCYRG---ICYR 14

RESULT 11
 JX0125
 tachyplesin III - horseshoe crab (Tachyplesus gigas)
 C:Species: Tachyplesus gigas
 C:Date: 30-Jun-1992 #sequence_revision 30-Jun-1992 #text_change 11-Jul-1997
 C:Accession: JX0125
 R:Muta, T.; Fujimoto, T.; Nakajima, H.; Iwanaga, S.
 J. Biochem. 108, 261-266, 1990
 A:Title: Tachyplesins isolated from hemocytes of southeast Asian horseshoe crabs (Car

ssing intermediate of its precursor.

A:Reference number: JX0124; MUID:91035357

A:Accession: JX0125

A:Molecule type: protein

A:Residues: 1-17 <MUT>

A:Experimental source: hemocyte

C:Keywords: amidated carboxyl end

F:3-16,7-12/Disulfide bonds: #status predicted

F:17/Modified site: amidated carboxyl end (Arg) #status experimental

Query Match 37.8%; Score 29.5; DB 2; Length 17;

Best Local Similarity 53.8%; Pred. No. 38;

Matches 7; Conservative 0; Mismatches 3; Indels 3; Gaps 1;

QY 1 RVCAGRGTCYR 13

Db 5 RVCYRG---ICYR 14

RESULT 12

A30068*

tachyplesin - horseshoe crab (Tachyplesus tridentatus)

C:Species: Tachyplesus tridentatus

C>Date: 28-Feb-1990 #sequence_revision 28-Feb-1990 #text_change 18-Jun-1993

C:Accession: A30068

R:Nakamura, T.; Furunaka, H.; Miyata, T.; Tokunaga, F.; Muta, T.; Iwanaga, S.; Niwa, M.;

J. Biol. Chem. 263, 16709-16713, 1988

A:Title: Tachyplesin, a class of antimicrobial peptide from the hemocytes of the horseshoe

A:Reference number: A30068; MUID:89034158

A:Accession: A30068

A:Molecule type: protein

A:Residues: 1-17 <NAK>

Query Match

Best Local Similarity 37.8%; Score 29.5; DB 2; Length 17;

Matches 7; Conservative 0; Mismatches 3; Indels 3; Gaps 1;

QY 1 RVCAGRGTCYR 13

Db 5 RVCYRG---ICYR 14

RESULT 13

JU0123

tachyplesin II - horseshoe crab (Tachyplesus tridentatus)

C:Species: Tachyplesus tridentatus

C>Date: 31-Mar-1990 #sequence_revision 31-Mar-1990 #text_change 11-Jul-1997

C:Accession: JU0123

R:Miyata, T.; Tokunaga, F.; Yoneya, T.; Yoshikawa, K.; Iwanaga, S.; Niwa, M.; Takao, T.;

J. Biochem. 106, 663-668, 1989

A:Title: Antimicrobial peptides, isolated from horseshoe crab hemocytes, tachyplesin II,

A:Reference number: A9194; MUID:90110066

A:Accession: JU0123

A:Molecule type: protein

A:Residues: 1-17 <MIY>

C:Comment: The peptide is one of the antimicrobial peptides found in the Japanese horseshoe

C:Keywords: amidated carboxyl end

F:3-16,7-12/Disulfide bonds: #status predicted

F:17/Modified site: amidated carboxyl end (Arg) #status experimental

Query Match

Best Local Similarity 37.8%; Score 29.5; DB 2; Length 17;

Matches 7; Conservative 0; Mismatches 3; Indels 3; Gaps 1;

QY 1 RVCAGRGTCYR 13

Db 5 RVCYRG---ICYR 14

RESULT 14

S59504

ferric pseudobactins receptor protein RF2 - Pseudomonas putida (fragment)

C:Species: Pseudomonas putida

C>Date: 20-Jul-1996 #sequence_revision 13-Mar-1997 #text_change 10-Jul-1998

C:Accession: S59504

R:Koster, M.; Ova, W.; Bitter, W.; Weisbeek, P.

Mol. Gen. Genet. 248, 735-743, 1995

A:Title: Multiple outer membrane receptors for uptake of ferric pseudobactins in pseu

A:Reference number: S59503

A:Accession: S59504

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-185 <ROS>

C:Superfamily: tonB-dependent receptor carboxyl-terminal homology

F:1-185/Domain: tonB-dependent receptor carboxyl-terminal homology (fragment) <TNC>

Query Match

Best Local Similarity 44.9%; Score 35; DB 2; Length 185;

Matches 5; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 RVCAGRGTCY 12

Db 30 KCGAGPASACY 41

RESULT 15

JX0124

tachyplesin I precursor - horseshoe crab (Carcinoscorpius rotundicauda)

C:Species: Carcinoscopus rotundicauda

C>Date: 30-Jun-1992 #sequence_revision 30-Jun-1992 #text_change 11-Jul-1997

C:Accession: JX0124

R:Muta, T.; Fujimoto, T.; Nakajima, H.; Iwanaga, S.

J. Biochem. 108, 261-266, 1990

A:Title: Tachyplesins isolated from hemocytes of southeast Asian horseshoe crabs (Car

ssing intermediate of its precursor.

A:Reference number: JX0124; MUID:91035357

A:Accession: JX0124

A:Molecule type: protein

A:Residues: 1-19 <MUT>

A:Experimental source: hemocyte

C:Keywords: amidated carboxyl end

F:1-17/Product: tachyplesin I #status experimental <MAT>

F:3-16,7-12/Disulfide bonds: #status predicted

F:17/Modified site: amidated carboxyl end (Arg) (amide in mature form from following

Query Match

Best Local Similarity 37.8%; Score 29.5; DB 2; Length 19;

Matches 7; Conservative 0; Mismatches 3; Indels 3; Gaps 1;

QY 1 RVCAGRGTCYR 13

Db 5 RVCYRG---ICYR 14

Search completed: May 4, 1999, 08:18:17

Job time: 6969 sec